Ageing & Society **25**, 2005, 111–130. © 2005 Cambridge University Press DOI: 10.1017/S0144686X04002740 Printed in the United Kingdom III

# Predicting transitions in the use of informal and professional care by older adults

# SANDRA W. GEERLINGS\*†, ANNE MARGRIET POT\*\*†, JOS W. R. TWISK†‡ and DORLY J. H. DEEG\*†

# ABSTRACT

To prepare the care system for a rising population of older people, it is important to understand what factors predict the use of care. This paper reports a study of transitions in the use of informal and professional care using Andersen-Newman models of the predictive predisposing, enabling and need factors. The study has drawn on Longitudinal Ageing Study Amsterdam (LASA) data on the use of care and the contextual factors. The data were collected at three-yearly intervals from a random, sex- and age-stratified, population-based sample of adults aged 55-85 vears. In summary, the findings for those who initially did not receive care were that almost one-third received some kind of care three years later, most of which was provided by informal care-givers. Need factors were important predictors of most transitions in care, and predisposing and enabling factors, such as age, partner status and income, also played a role. On the relationship between informal and professional care, evidence was found for both 'compensatory processes', i.e. informal care substitutes for professional care, and 'bridging processes', *i.e.* informal care facilitates professional care. In view of the increasing discrepancy between the demand for professional care and its supply, the significant impact of predisposing and enabling factors offers opportunities for intervention.

KEY WORDS - need for care, care transitions, professional care, subsidised care.

#### Introduction

As for most developed nations, the current demographic projections for The Netherlands indicate that approximately a quarter of the population will be 65 or more years of age by 2040 (Central Bureau of Statistics 1997).

<sup>\*</sup> Department of Psychiatry, VU University Medical Center, Amsterdam.

<sup>†</sup> Institute for Research in Extramural Medicine, VU University Medical Centre, Amsterdam.

<sup>\*\*</sup> Department of Nursing Home Medicine, VU University Medical Centre, Amsterdam.

<sup>&</sup>lt;sup>‡</sup> Department of Clinical Epidemiology and Biostatistics, VU University Medical Centre, Amsterdam.

The population at the oldest ages will increase most rapidly, and even if age-specific morbidity and dependency rates moderate, the requirement for long-term care will continue to increase. In order to prepare the care system, it is of great importance to understand what factors predict the use of care. Most studies on the predictors of care utilisation rely on cross-sectional data and therefore describe differences between users and non-users. To explain the use of care, however, it may be more useful to focus on the process of becoming a user. To construct a theory of the factors that influence the use of care, a longitudinal design is required, in which 'transitions in care' are the dependent or outcome variables; this enables the changing situation of the older person to be taken into account (Crets 1996).

It is also important in the study of care transitions to consider simultaneously informal and professional care. Informal care includes support from partners, adult children and other relatives, friends and neighbours, whereas professional care comprises institutional care and home-care provided by subsidised or fee-paid professionals. Theorists have not agreed on how to link the relationship between informal and professional care (Logan and Spitze 1994; Lyons and Zarit 1999). In the 'hierarchical compensatory model' (Cantor 1979), professional services are considered to be a substitute for informal care when the latter are not forthcoming, and informal support is also represented as a bridge between the older person and professional services, bringing the person into contact with professional service providers and thus facilitating professional care (George 1987; Sussman 1976).

To predict the use of care, Andersen and Newman's (1973) behavioural model of health services use has often been employed (Andersen 1995; Borrayo *et al.* 2002; Kempen and Suurmeijer 1991). It distinguishes 'predisposing', 'enabling' and 'need' factors. Predisposing factors reflect the propensity to use services independent of the personal circumstances and experiences that may cause the need for care, and typically are sociodemographic variables. Enabling characteristics relate to the variable care resources available to the individual, as exemplified by partner status (the availability of a partner), the level of urbanisation (the availability of formal services), and the availability of informal care. Need factors pertain to the needs of the person and represent the most immediate stimulus to obtain care: they include physical and mental health status.

The Longitudinal Ageing Study Amsterdam (LASA) has created good data resources with which to study care transitions among older people. First, the longitudinal population-based design with three-yearly data collection intervals permits the prediction of the 'onset' of care among initial nonusers. Second, the presence of data on both informal and professional care enables close examination of the relationships between the two sources. Third, information is available on the three types of influences in Andersen and Newman's model, and finally, the longitudinal design makes it possible to estimate the effects of changes over time in the predictor variables (particularly the need factors). The study reported in this paper had two aims: first, to describe transitions in the use of care, with a focus on the transitions from 'no care' to 'informal home-care', and from either 'no care' or 'only informal home-care' to 'professional home-care' or 'institutional care'; and secondly, to assess the explanatory value of (changes in) predisposing, enabling and need factors.

# Methods

#### Sampling and procedures

LASA is an inter-disciplinary study of the predictors and consequences of changes in autonomy and wellbeing among older people (full details of the sampling procedures and response rates have been described elsewhere: Deeg and Westendorp-de Serière 1994; Deeg et al. 1998). The main points are that a representative random sample of older adults (aged 55-85 years), stratified for age and sex, was drawn from the population registries of 11 municipalities in three regions of The Netherlands and used in two studies. First, 3,805 respondents were interviewed for the NESTOR project on 'Living arrangements and social networks of older adults' (NESTOR-LSN) (response rate 62.3%) (Broese van Groenou et al. 1995). About ten months later, 3,107 (81.7%) of the original respondents took part in the LASA baseline interview.<sup>1</sup> Follow-up measurements using exactly the same instruments and procedures took place at three-yearly intervals (Smit, De Vries and Poppelaars 1998). This study has analysed data from the baseline  $(t_0)$  interview in 1992/93, from the first follow-up interview  $(t_1)$ in 1995/96, and the second follow-up interview  $(t_2)$  in 1998/99.

#### Measurements

As the *dependent variable*, the use of care was measured by asking the respondents whether they received help for instrumental activities of daily living, such as cleaning the house and shopping (household care), and if so, from which sources. These questions were also asked about help received for the basic activities of daily living, such as bathing and dressing (personal care). Informal, subsidised and private home or domiciliary care, and institutional care were all distinguished. Informal home-care is unpaid, non-organised help offered by partners, adult children and other relatives, friends and neighbours. A distinction was made between subsidised and private home-care because the former is provided by professional non-profit organisations and the government subsidises most of the costs. Governmental committees organise the allocation and decide on the eligibility of disabled older people for subsidised care. Private homecare is provided by for-profit organisations. It has a growing role in the support of older people (Portrait, Lindeboom and Deeg 2000). The bulk of institutional care is provided in subsidised residential and nursing homes (geriatric units in general and psychiatric hospitals provide the remainder).

#### Independent variables

The independent variables were divided into predisposing, enabling and need factors (Anderson and Newman 1973). *Predisposing factors* included sex, age and education. Education was measured on an eight-point ordinal scale that ranged from 'uncompleted primary school' to 'completed university education'. For the analysis reported in this paper, education was dichotomised as 'low' (no more than completed primary school, ages 6 to 12 years), and 'middle to high' (some secondary education up to and including university degree). *Enabling variables* were (changes in) partner status, level of urbanisation, income and the presence of informal homecare. *Need variables* included (changes in) chronic physical diseases, functional limitations, depression, cognitive impairment, and self-rated health.

The presence of chronic physical diseases was assessed by asking the participants whether they had any of the following diseases: cardiac disease, peripheral atherosclerosis of the abdominal aorta or the arteries of the lower limb, stroke, diabetes mellitus, lung disease (asthma or chronic obstructive pulmonary disease), cancer or arthritis (rheumatoid arthritis or osteoarthritis) (Kriegsman et al. 1996). Based on two consecutive measurements, respondents were divided into four categories: 'no diseases at successive measurements' (the reference category), 'a constant number (at least one) of chronic diseases at successive measurements', 'incident comorbid disease(s)', and 'incident first disease(s)'. Functional limitations were assessed by the following activities: climbing up and down a staircase of 15 steps without stopping, cutting one's own toenails, and using one's own or public transport (Kriegsman et al. 1997; Van Sonsbeek 1988). The response categories were 'yes, without difficulty', 'yes, with difficulty', 'only with help' and 'no, I cannot'. For this study, the presence of functional limitations was dichotomised as follows: 'no difficulty with any functional limitations' and 'have difficulty with one or more functional limitations'. Based on two consecutive measurements, respondents were divided into four categories: 'no functional limitations at successive measurements'

(the reference category), 'functional limitations at successive measurements' (has limitations, stable), 'functional limitations at  $t_x$  but not at  $t_{x+1}$ ' (recovery from limitations), and 'functional limitations at  $t_{x+1}$  but not at  $t_x$ ' (incidence of limitations).

Depressive symptoms were measured using the Center for Epidemiologic Studies Depression Scale (CES-D) (Radloff 1977), a 20-item self-report scale (range 'o' to '60') designed to measure depressive symptomatology in the general population. To identify clinically relevant depressive syndromes, the generally accepted threshold of '16' was applied (Berkman et al. 1986). A clinically relevant change was defined as a decrease or an increase of at least five points on the CES-D that crossed the threshold of '16' (Jacobson and Truax 1991). Based on two consecutive measurements, respondents were divided into four categories: 'no depression at successive measurements' (the reference category), 'depression at successive measurements' (chronic depression), 'depression at  $t_x$  but not at  $t_{x+1}$ ' (remission), and 'depression at  $t_{x+1}$  but not at  $t_x$ ' (incidence of depression) (Geerlings, S. W. et al. 2000). Cognitive impairment was measured using the Mini-Mental State Examination (MMSE) which has a range from 'o' to '30' and a threshold score of '24' (Folstein, Folstein and McHugh 1975; Tombaugh and McIntyre 1992). In line with previous research (Geerlings, M. I. et al. 2000), a critical change of the MMSE score was defined as a decrease of more than three points that crosses the threshold. Based on two consecutive measurements, respondents were divided into three categories: 'no cognitive impairment at successive measurements' (the reference category), 'cognitive impairment at successive measurements' (impaired, stable), and 'cognitive impairment at  $t_{x+1}$  but not at  $t_x$ ' (incidence of impairment). Self-rated health was measured by asking respondents to evaluate their own health, and the response categories were 'excellent', 'good', 'fair', 'sometimes good/bad' and 'poor' (Central Bureau of Statistics 1989). For this study, self-rated health was dichotomised as follows: 'excellent or good' and 'fair to poor'; and change in self-rated health over two consecutive measurements into four categories: 'excellent or good at both' (the reference category), 'fair to poor' at both, 'fair to poor at  $t_x$  and excellent or good at  $t_{x+1}$  (improvement), and 'excellent or good at  $t_x$  and fair to poor at  $t_{x+1}$ ' (decline).

#### Data analysis

Data were analysed in four stages. First, attrition to follow-up was studied using chi-squared tests and logistic regression. Secondly, transitions in care between  $t_0$  and  $t_1$  and between  $t_1$  and  $t_2$  were described and differences in

proportions tested by calculating confidence intervals. Thirdly, care transitions were predicted for respondents with valid data at  $t_0$  and  $t_1$ . For the 983 respondents without care at  $t_0$ , logistic regression models were used to predict the transition to informal home-care at  $t_1$ . In the same way, for the 1,718 respondents with no care or informal home-care at t<sub>0</sub>, multinomial regression models were used to predict the transition at t<sub>1</sub> to professional care (subsidised home-care, private home-care and institutional care) – informal home-care at  $t_0$  was then an independent variable. Odds ratios and 95 per cent confidence intervals were calculated for the bivariate associations between the predisposing, enabling and need variables and the care transitions. The predisposing, enabling or need variables that were shown by the bivariate analyses to be significant predictors were then entered into regression models of the care transitions. Finally, other variables that were shown as significant predictors in the former analyses were entered into the final model that included all dimensions of the Andersen and Newman model (1973). Since the study sample was stratified for sex and age, these variables were entered in all models regardless of their significance. In the final stage of the analysis, to validate the models, all analyses were repeated for respondents with valid data at t<sub>1</sub> and at t<sub>2</sub>.

# Results

#### Sample characteristics and attrition analyses

The characteristics of the baseline or  $t_0$  sample of 3,107 respondents are presented in Table 1. The relatively high proportions of men (49.5%) and of older-old adults (mean age 70.8 years) reflect the stratified sampling design. As a result of over-sampling of the more vulnerable groups, (physical) health problems are well represented in the baseline sample. A majority of the respondents at baseline received some kind of care: 84.9 per cent received only household care, one per cent received only personal care, and 14.1 per cent received both household and personal care. Since combinations of informal care and professional care were relatively rare (n = 172), and of these 49 per cent received both informal and professional home-care, these were categorised as receiving professional care only. Informal home-care was provided mostly by partners (69.2%).

Data on transitions in care between  $t_0$  and  $t_1$  and the required predictor variables were available from 2,259 respondents.<sup>2</sup> Attrition due to mortality was associated with nearly all baseline characteristics (with p values of <0.01 or less): higher age, males, low education, not (or no longer) having a partner, highly urbanised localities, one or more chronic physical

Attribute	Percentage	Attribute	Percentage
Sex :		Depression:	
Female	51.5	No	84.0
Education ·		Yes	14.4
Low	55.5	Unknown	1.6
Unknown	0.3	Cognitive impairment:	
Partner status.	5	No	88.5
Partner	66 <del>-</del>	Yes	11.0
No partner	00.5	Unknown	0.5
Unknown	0.2	Self-rated health:	
I Tak anisation .		Excellent or good	62.0
Utoanisation: High	o <del>7</del> 6	Fair to poor	37.2
Tingii	27.0	Unknown	0.8
Income:	2	Care use.	
High	26.1	No care	46.7
Chronic physical	diseases :	Informal home-care	40.7 25.4
No	25.7	Subsidized home-care	-0.4
Yes	73.7	Private home-care	13.1
Unknown	0.6	Institution	4.1
Functional limita	tions:	Unknown	0.7
No	57.8	Average age	70 8 yrs <sup>1</sup>
Yes	40.8	Sample size	2 107
Unknown	- 1.4	Sumpre Size	3,10/

TABLEI. Characteristics of the baseline sample

Note: 1. Standard deviation 8.8 years.

diseases, one or more functional limitations, impaired cognition, fair to poor self-rated health, and the use of care at  $t_0$ . Attrition through reasons other than mortality was associated at  $t_0$  with higher age, low education, highly urbanised localities, one or more functional limitations, impaired cognition, depression, and the use of care, but was *not* associated with sex, not (or no longer) having a partner, higher income, one or more chronic physical diseases, and fair to poor self-rated health. Data on transitions in care between  $t_1$  and  $t_2$  and the required predictor variables were available for 1,780 respondents.<sup>3</sup> Essentially the same associations between attrition and baseline characteristics were found as for transitions from  $t_0$  to  $t_1$ .

#### Transitions in care

The transitions in care between  $t_0$  and  $t_1$  and between  $t_1$  and  $t_2$  are presented in Table 2. A separate category has been included for those respondents who died before the follow-up. The upper section of Table 2 shows that among the initial non-users, 27.1 per cent received some kind

			Des	stination car	e-type		
Origin care-type	No care	Informal home-care	Subsidised home-care	Private home-care	Institutional care	Deceased	Total
At t <sub>0</sub>			Perce	ntages by colu	mn at t <sub>1</sub>		
No care	68.6	14.2	4.I	8.3	0.5	4.2	(48.0)
Informal home-care	42.8	34.9	6.5	5.7	2.7	7.3	(25.4)
Subsidised home-care	4.5	2.7	55.9	5.5	10.0	21.4	(8.9)
Private home-care	9.0	5.8	6.7	66.6	2.9	9.0	(13.9)
Institutional care	0.0	0.0	I.I	0.0	59.3	39.6	(3.7)
Total (sample size)	1124	414	<sup>2</sup> 37	375	109	210	2,469
Total (column %)	45.5	16.8	9.6	15.2	4.4	8.5	100
At t <sub>1</sub>			Perce	entages by colu	mn at $t_2$		
No care	64.2	17.9	6.3	6.1	0.9	4.7	(51.9)
Informal home-care	29.0	45.8	8.7	8.4	2.6	5.5	(17.9)
Subsidised home-care	4.3	2.2	62.0	2.7	9.8	19.0	(9.5)
Private home-care	10.5	4.6	7.7	68.o	3.1	6.2	(16.8)
Institutional care	-	_	1.3	-	58.7	40.0	(3.9)
Total (sample size)	785	356	233	316	90	151	1,931
Total (column %)	40.7	18.4	12.1	16.4	4.7	7.8	100

T A B L E 2. Transitions in the use of care and mortality between  $t_0$  and  $t_1$  and between  $t_1$  and  $t_2$ 

Note: Figures in parentheses are row percentages.

of care three years later. More than half of them (14.2%) of the total) received informal home-care. The instability of informal home-care over time is striking: only 34.9 per cent received informal home-care at both  $t_0$  and  $t_1$ . There was however more stability in the receipt of the various types of professional care, for between 55.9 and 66.6 per cent were clients at both dates. The instability of informal home-care was mainly the result of a large number (42.8%) losing this support over time. Only a few respondents reported the transition from professional care to informal care.

Compared to the initial non-users, the respondents who received informal home-care or professional care were more likely to die between  $t_0$  and  $t_1$  (p = < 0.01). Mortality was highest among the respondents in institutional care (39.6%), followed by those who received subsidised home-care (21.4%). The transitions between  $t_1$  and  $t_2$  are set out in the lower section of Table 2. They resembled those during the earlier period except that the transitions from 'no care' to 'informal home-care' or 'professional care' were more frequent (31.2% versus 27.1%, p = < 0.01), and

Variables	OR (95 $\%$ CI) $^{1,2}$	Variables	OR (95 % CI) 1,2	
Need variables		Predisposing variables		
Chronic physical diseases ≥ 1 diseases, stable Incident co-morbid disease(s) Incident first disease(s)	1.30 (0.86–1.96) 1.90 (1.13–3.17) 2.15 (1.31–3.51)	Age Sex Female	<b>1.04 (1.02–1.07)</b> 0.77 (0.55–1.08)	
Functional limitations Limited, stable Recovery Incident limitations	<b>2.00 (1.32–3.03)</b> 1.15 (0.54–2.42) 1.19 (0.76–1.87)	Education Low	1.55 (1.11–2.16)	
Depression Chronic depression Remission Incident depression	0.44 (0.13–1.46) 0.81 (0.37–1.74) 1.04 (0.54–1.99)	<b>Enabling variables</b> Partner status No partner Lost partner t <sub>0</sub> –t <sub>1</sub>	s 0.57 (0.37–0.87) 0.72 (0.33–1.57)	
Cognitive impairment Impaired, stable Incident impairment	1.75 (0.89–3.47) 0.93 (0.43–2.01)	Urbanisation High	0.38 (0.25–0.56)	
Self-rated health Fair to poor, stable Improvement Decline	<b>1.91 (1.27–2.87)</b> 1.46 (0.87–2.43) 1.34 (0.76–2.34)	Income High <b>Sample size</b>	1.32 (0.92–1.90) 983	

T A B L E 3. Bivariate associations between predisposing, enabling and need variables and the transition from no care at  $t_0$  to informal home-care at  $t_1$ 

Notes: 1. OR, odds ratio; CI, confidence interval. 2. Significant odds ratios are in bold.

there was greater continuity among those who received informal homecare (45.8% versus 34.9%, p = < 0.001).

#### Predicting transitions in care

# Transition from no care to informal care

Table 3 displays the bivariate associations between the predisposing, enabling and need variables and the transition from 'no care' at  $t_0$  to 'informal home-care' at  $t_1$ . The significantly predictive factors were: among the *predisposing variables*, higher age and low education; among the *enabling variables*, having no partner and rural (little urbanised) localities; and among the *need variables*, the incidence of first and co-morbid diseases, stable functional limitations, and stable 'fair to poor' self-rated health. Additional analyses were performed to differentiate the relationships between seven chronic physical diseases and informal care. The transition to informal care at  $t_1$  was predicted by the prevalence at  $t_0$  of diabetes mellitus (odds ratio (OR)=2.05, 95% confidence interval (c.i.) 1.02–4.15) and stroke (OR = 3.47, c.i. 1.50–8.04). Arthritis, cardiac disease, lung disease, cancer

Independent variables	OR $(95\% \text{ CI})^{1,2}$	OR (95 % CI) <sup>1,2</sup>	
Predisposing variables	Model IA <sup>3</sup>	Model II <sup>3</sup>	
Age	1.04 (1.01–1.06)	1.05 (1.02–1.07)	
Education			
Low	1.55 (1.09–2.21)		
Enabling variables	Model IB <sup>3</sup>		
Partner status			
No partner	0.59 (0.39–0.91)	0.50 (0.30–0.81)	
Lost partner $t_0-t_1$	0.71 (0.33–1.56)	0.54 (0.23–1.19)	
Urbanisation			
High	0.38 (0.26–0.57)	0.31 (0.20-0.48)	
Need variables	Model IC <sup>3</sup>		
Chronic physical diseases			
≥1 diseases, stable	1.01 (0.64–1.60)	1.09 (0.70–1.72)	
Incident co-morbid disease(s)	1.35 (0.76–2.40)	1.32 (0.74–2.35)	
Incident first disease(s)	1.86 (1.12–3.12)	2.01 (1.19–3.41)	
Functional limitations			
Limited, stable	1.64 (1.02–2.64)	1.73 (1.05–2.86)	
Recovery	0.91 (0.41-2.04)	1.17 (0.53–2.61)	
Incident limitations	1.06 (0.67–1.69)	1.07 (0.66–1.75)	

T A B L E 4. Multivariate associations between predisposing, enabling and need variables and the transition from no care at  $t_0$  to informal home-care at  $t_1$ 

*Notes*: I. OR, odds ratio; CI, Confidence interval. 2. Results are printed for significant variables only and significant odds ratio's are in bold. 3. Model IA, only predisposing variables; Model IB, only enabling variables; Model IC, only need variables. Model II, model including predisposing, enabling and need variables. The sample size for all models was 983.

and peripheral arteriosclerosis were *not* associated with the transition to informal home-care.

Table 4 shows the multivariate associations between the predisposing, enabling and need variables and the transition from no care at to informal home-care at  $t_1$ . The results from the first two models resemble those from the bivariate analyses. In the model that included only predisposing variables (IA), both higher age and low education were predictive. In the model that included only enabling variables (IB), having no partner as well as the locality being little urbanised were significantly associated. The model of need variables (IC) identified the incidence of first chronic physical diseases and stable functional limitations as predictive while, contrary to the bivariate analyses, the incidence of co-morbid diseases and stable 'fair to poor' selfrated health were insignificant. All statistically significant predictors in these three models were entered into the final model (II), which found that the transition was best predicted by higher age, having no partner, the locality being little urbanised, the incidence of first chronic physical diseases, and stable functional limitations. To validate the models, the analyses were repeated for the respondents with valid data at  $t_1$  and  $t_2$ . The results

were very similar, but the transition to informal home-care was no longer predicted by higher age (OR = 0.99, c.i. 0.97–1.02), while the incidence of functional limitations (OR = 1.68, c.i. 1.06-2.67) and a decline in self-rated health (OR = 1.65, c.i. 1.04-2.65) entered as significant predictors.

# The transition from 'no care' or 'informal care' to 'professional care'

Table 5 shows the bivariate results of the multinomial regression analyses of the transition between  $t_0$  and  $t_1$  from 'no care' or 'informal home-care' to 'professional care' (subsidised home-care, private home-care or institutional care). From the predisposing variables, higher age was predictive for all possible transitions, while low education predicted the transitions to subsidised home-care and institutional care, and high education predicted the transition to privately purchased home-care. The transition to subsidised home-care was predicted by almost all enabling variables: having no partner or losing a partner, a highly urbanised locality, and informal home-care at t<sub>0</sub>. The transitions to private home-care and institutional care were both predicted by having no partner. Furthermore, an association was found between higher income and the transition to privately purchased home-care and between informal homecare at  $t_0$  and institutional care at  $t_1$ . All 'disabling' need variables were associated with receiving subsidised home-care at  $t_1$ , but conversely, the remission of depression, recovery from functional limitations and improvement in self-rated health were insignificant. The transition to privately purchased home-care was predicted only by chronic physical diseases (whether stable or incident co-morbid diseases). Stable functional limitations, incident functional limitations, incident depression and stable or incident cognitive impairment were associated with the transition to institutional care.<sup>4</sup>

The results of the multivariate multinomial regression analyses are presented in Table 6. Because of the few (23) respondents, those in institutional care were excluded. In Model IA (predisposing variables only), higher age was associated with the transition to subsidised home-care as well as the transition to private home-care. The transition to private home-care was also predicted by being female and having high education. Model IB (enabling variables only) shows that having no partner or losing a partner and being in receipt of informal home-care at t<sub>0</sub> predicted the transition to subsidised home-care, and that having no partner and higher income predicted the transition to private home-care. In Model IC (need variables only), the transition to subsidised home-care was predicted by the onset of co-morbid chronic physical diseases, stable functional limitations, stable cognitive impairment and a decline in self-rated health. In

Independent variables	Subsidised home-care OR $(95\% \text{ CI})^{1,2}$	Private home-care OR $(95 \% \text{ CI})^{1,2}$	Institutional care OR $(95\% \text{ CI})^{1,2}$	
Predisposing variables Age	1.13 (1.09–1.16)	1.06 (1.04–1.09)	1.29 (1.19–1.41)	
Sex Female	1.34 (0.87–2.05)	1.15 (0.81–1.64)	1.91 (0.81–4.54)	
Education Low	1.75 (1.14–2.69)	0.52 (0.35–0.77)	3.86 (1.51-9.85)	
Enabling variables Partner status No partner Lost partner t <sub>0</sub> -t <sub>1</sub>	3.09 (1.95–4.90) 3.88 (1.96–7.68)	<b>1.78 (1.22–2.62)</b> 1.10 (0.49–2.47)	11.05 (4.05-30.20) 2.59 (0.30-22.29)	
Urbanisation High	1.55 (1.01-2.39)	1.12 (0.78–1.62)	0.85 (0.35-2.08)	
Income High	0.66 (0.39–1.14)	1.64 (1.14–2.30)	0.79 (0.29–2.15)	
$\begin{array}{c} Informal \ home-care \ t_0 \\ Yes \end{array}$	1.69 (1.10-2.59)	0.74 (0.50–1.10)	5.71 (2.34–14.57)	
Need variables Chronic physical diseases ≥1 diseases, stable Incident co-morbid disease(s) Incident first disease(s)	3.78 (1.87–7.65) 7.82 (3.71–16.47) 3.62 (1.58–8.28)	<b>2.04</b> ( <b>1.30–3.19</b> ) <b>2.70</b> ( <b>1.57–4.65</b> ) 1.03 (0.52–2.06)	1.11 (0.40–3.08) 2.07 (0.65–6.60) 1.11 (0.28–4.33)	
Functional limitations Limited, stable Recovery Incident limitations	<b>9.90</b> (5.43–18.05) 1.43 (0.32–6.41) <b>2.97</b> (1.40–6.32)	1.45 (0.95–2.21) 0.30 (0.07–1.26) 1.17 (0.73–1.88)	<b>32.75</b> ( <b>4.27–251.39</b> ) 10.01 (0.62–161.63) <b>14.87</b> ( <b>1.73–127.84</b> )	
Depression Chronic depression Remission Incident depression	<b>2.36 (1.03–5.39)</b> 0.97 (0.35–2.75) <b>3.24 (1.80–5.83)</b>	0.36 (0.09–1.47) 0.77 (0.33–1.80) 1.28 (0.68–2.40)	2.84 (0.63–12.78) 1.14 (0.41–9.88) 3.41 (1.10–10.55)	
Cognitive impairment Impaired, stable Incident impairment	3.88 (2.12–7.11) 2.80 (1.38–5.68)	0.82 (0.35–1.92) 0.74 (0.29–1.87)	23.54 (7.30–75.83) 36.42 (12.13–109.34)	
Self-rated health Fair to poor, stable Improvement Decline	<b>3.71 (1.99–6.93)</b> 1.84 (0.87–3.88) <b>3.96 (2.35–6.67)</b>	1.17 (0.74–1.84) 0.81 (0.43–1.52) 1.58 (0.95–2.65)	2.00 (0.76–5.31) 0.96 (0.21–4.40) 2.14 (0.66–6.92)	

T A B L E 5. Bivariate associations between predisposing, enabling and need variables and the transition from no care or informal care at  $t_0$  to subsidised home-care, private home-care or institutional care at  $t_1$ 

Notes: 1. OR, odds ratio; CI, confidence interval. 2. Significant odds ratios are in bold. Sample size in all models is 1,718.

addition, stable chronic diseases and incident co-morbid diseases predicted the transition to privately purchased home-care.

The final model (II) included all the statistically significant variables in the first three models, among which the following predicted the transition

	Subsidised	home-care	Private home-care		
Independent variables	OR (95 % CI) <sup>1,2</sup>	OR (95 % CI) <sup>1,2</sup>	OR (95% CI) <sup>1,2</sup>	OR (95 % CI) <sup>1,2</sup>	
Predisposing variables	Model IA <sup>3</sup>	Model II <sup>3</sup>	Model IA <sup>3</sup>	Model II <sup>3</sup>	
Age	1.13 (1.09–1.16)	1.08 (1.05-1.12)	1.08 (1.05–1.10)	1.08 (1.05–1.11)	
Sex Female	-	-	1.59 (1.09-2.32)	_	
Education Low	-	-	0.37 (0.24–0.56)	0.39 (0.25–0.61)	
Enabling variables Partner status	Model IB <sup>3</sup>		Model IB <sup>3</sup>		
No partner Lost partner t <sub>0</sub> —t <sub>1</sub>	3.24 (2.03–5.18) 3.97 (2.00–7.92)	1.85 (1.04–3.32) 3.14 (1.42–6.95)	<b>1.78 (1.21–2.63)</b> 1.10 (0.49–2.45)	_	
Income High	_	_	1.81 (1.23–2.68)	1.82 (1.23–2.67)	
$\begin{array}{c} \mbox{Informal home-care } t_0 \\ \mbox{Yes} \end{array}$	1.99 (1.28–3.09)	_	_	_	
Need variables Chronic physical diseases	Model IC <sup>3</sup>		Model IC <sup>3</sup>		
≥1 diseases, stable Incident co-morbid disease(s)	2.26 (0.98–5.20) <b>3.68 (1.51–8.98</b> )	2.09 (0.90–4.83) 4.00 (1.64–9.79)	2.49 (1.51–4.10) 3.10 (1.71–5.64)	2.32 (1.41–3.82) 2.75 (1.50–5.03)	
Incident first disease(s)	2.41 (0.94–6.18)	1.96 (0.75–5.12)	1.10 (0.54–2.25)	1.04 (0.51–2.14)	
Functional limitations Limited, stable Recovery Incident limitations	<b>5.30 (2.73–10.29)</b> 1.06 (0.23–4.87) 1.89 (0.86–4.17)	<b>2.77 (1.37–5.62)</b> 0.81 (0.17–3.78) 1.26 (0.56–2.84)		0.78 (0.46–1.32) <b>0.20 (0.05–0.86)</b> 0.78 (0.46–1.32)	
Cognitive impairment Impaired, stable Incident impairment	<b>2.46 (1.20–5.03)</b> 1.90 (0.84–4.30)	_	_		
Self-rated health Fair to poor, stable Improvement Decline	1.53 (0.80–2.90) 1.26 (0.56–2.84) <b>2.15 (1.07–4.32)</b>	<b>2.11 (1.04–4.19)</b> 1.51 (0.67–3.43) <b>2.53 (1.26–5.07)</b>			

T A B L E 6. Multivariate associations between predisposing, enabling and need variables and the transition from no care or informal care at  $t_0$  to subsidised home-care or private home-care at  $t_1$ 

*Notes*: I. OR, odds ratio; CI, Confidence interval. 2. Results are presented for significant variables only and significant odds ratios are in bold. 3. Model IA, only predisposing variables; Model IB, only enabling variables; Model IC, only need variables. Model II, model including predisposing, enabling and need variables. Sample size in all models is 1,695.

to subsidised home-care at  $t_1$ : higher age, having no partner or losing a partner, incident co-morbid diseases, stable functional limitations, stable 'fair to poor' self-rated health, and a decline in self-rated health. Closer inspection of the model coefficients showed that the association between informal home-care at t<sub>0</sub> and, after inclusion of the need variables, the transition to subsidised home-care was no longer statistically significant, and that the odds ratio reduced to 1.39 (p=0.18). The transition to privatelypurchased home-care was associated with higher age, high education, high income, stable chronic diseases and incident co-morbid diseases. Recovery from functional limitations was negatively associated with the transition to private home-care. To validate the models, the analyses were repeated for the respondents with valid data at t<sub>1</sub> and t<sub>2</sub>. There was no change in the included predisposing variables, but for the enabling variables, an additional association (p=0.08) was found between informal home-care at t<sub>1</sub> and private home-care at t<sub>2</sub> (OR = 1.52, c.i. 0.95–2.41). The included need variables also changed in the model of the transition to privately purchased home-care.<sup>5</sup>

#### Discussion

This study has modelled transitions over two successive three-year intervals during the 1990s in the use of informal and professional care in a representative sample of Dutch older people, and has examined the influence of both the starting values of and changes in various predisposing, enabling and need factors. Various transitions between no care, informal home-care, subsidised home-care, privately purchased home-care and institutional care were observed. It has been shown that among those who were initially non-users, 27 to 31 per cent used some kind of care three years later. Consistent with much previous research in other developed nations, most of the care was provided by informal care-givers (Roe et al. 2001; Tennstedt et al. 1990), but unlike professional care, its receipt showed little stability over time, mostly because a sizable minority experienced the transition from 'informal home-care' to 'no care'. Additional analyses (not reported here) have shown that this retraction occurred most often among people with relatively favourable predisposing, enabling and need factors. Jette, Tennstedt and Branch's (1992) longitudinal study found informal care to be much more stable than in the LASA sample, which may be explained by the higher average age and greater frailty of their sample - they would have been more dependent upon prolonged care. Mortality within three years was associated with the use of care, and was highest among the respondents in institutional settings followed by those receiving subsidised home-care.

On the prediction of the transition to informal home-care, both the immediate need for care, associated with chronic physical diseases and functional limitations and certain predisposing and enabling factors (such as advanced age and having no partner), were shown to be necessary conditions. Turning to the transition from 'no' or 'informal' home-care to 'professional' care, the findings revealed the importance of distinguishing various sources of such care. There were differential effects of (changes in) the predisposing, enabling and need factors for subsidised and private home-care and for institutional care. Some variables (such as age) consistently affected all measures of care, while others (such as education) affected one measure in one direction and another in the opposite direction, and some (such as sex) had no effect at all. The need factors were especially important in predicting subsidised home-care and institutional care, while higher education and higher income were strong predictors of the transition to private home-care.

The finding that need factors were strongly associated with the transitions both to subsidised home-care and to institutional care was strengthened by the evidence that these sources of care positively associated with mortality. These interlinked associations support the notion that where professional care is provided, it is needed, which evinces the effectiveness of the governmental need-assessment committees that allocate subsidised care. Given the significant influence of predisposing and enabling factors, it cannot further be concluded, however, that care is provided to *all* those who need it. Several explanations have been offered for not receiving professional care given a demonstrable need: some people may have been declined care by the assessment committee, others may have been on a waiting list because of supply constraints, and others may not have asked for it (Portrait, Lindeboom and Deeg 2000). 'Unexpressed demand' may arise through unfamiliarity with care services or negative attitudes and beliefs about the quality of professional care (MaloneBeach, Zarit and Spore 1992; Zarit, Gaugler and Jarrott 1999); or because informal care-givers withhold an application to prevent feelings of guilt (Pot, Deeg and Knipscheer 2001); or because it is impossible for need-assessment committees repeatedly to re-assess people's care needs - mental health status, for instance, can be highly variable over time (Geerling, S. W. et al. 2000).

It was broadly found that informal and professional care operated independently, not complementarily, which supports the proposition that their receipt is dependent upon the tasks involved. Professional service providers may be less suited to unskilled home-care, for which the motivation of a partner or other close relative is especially advantageous. For other care tasks, particularly those which require specialised training or facilities, professional services may be more appropriate (Litwak 1985). Combinations of types of care generally reflect multiple needs. In the last phase of life, combinations of informal and professional care have been found to be rather common, especially for those with more severe and protracted illness (Klinkenberg 2004).

Support was found for both the 'hierarchical compensatory model' and the 'bridging hypothesis'. In general, older people with a partner were less likely to receive professional care, which is in line both with the view that informal care substitutes for professional services and with the hierarchical compensatory model. Conversely, the respondents who initially had informal home-care, three years later were more likely than others to receive professional care (particularly subsidised home-care and institutional care) because of their relatively poor health status. This finding supports the contention that informal care brings the person into contact with professional service providers and facilitates rather than substitutes for professional care, in line with the bridging hypothesis. Probably, the closeness or strength of an informal tie determines whether informal care has substitutive or bridging effects. It has been argued that the effect of informal care depends on the composition of the network: close relatives tend simply to substitute their own help for professional services, whereas friends and neighbours are more apt to perform complementary tasks (Logan and Spitze 1994; Ward, Sherman and LaGory 1984). Additional analyses (results not shown) showed a further differentiation of the relationship between informal care and professional care: the complementary relationship was clearest when non-partners were involved. As found in previous research (Tennstedt, Harrow and Sullivan 1996), the transition from professional care to informal care occurred only seldom and was therefore impossible to analyse statistically.

The transitions and their determinants for the first and second time intervals were quite similar (demonstrating the robustness of the models) but some differences were found. Stability in receiving informal homecare was noticeably higher during the second period, which might be partly explained by the need for care to become more intense, and by the strengthened motivation, skills and support of informal care providers (Hanson *et al.* 2002). Furthermore, the need factors became more important in predicting the transition to private home-care, which may be explained by changes in the availability, allocation and supply of subsidised care services in The Netherlands during the study period. In general, the reduction of the number of beds in hospitals and residential homes has increased the pressure on professional home-care services and resulted in longer waiting lists (Prismant 2000). Such supply restrictions may have necessitated the increased use of private resources.

Some limitations of the present study have to be acknowledged. The over-sampling of men and older-old subjects produced considerable

non-response through mortality and a higher attrition rate in the subjects with the worst health. As attrition was associated with the use of care as well as practically all predisposing, enabling and need factors, the strength of the associations between predictor variables and transitions in care is likely to have been under-estimated. A second limitation may be the relatively long period of three years between the measurements. Although stability was high for most sources of care, shorter intervals between the measurements are recommended in future studies, especially those pertaining to informal care, which was most variable. A third concern may be that chronic physical diseases and functional limitations were measured by selfreport. In a validation study, however, respondents' self-reports on chronic diseases were compared to information obtained from their GPs and proved reliable (Kriegsman et al. 1996). With respect to functional limitations, more objective measures such as performance tests in a more experimental setting have not been found to be superior to self-reports. Considerable discordance has even been reported between performance in a test and actual functioning at home (Glass 1998). Finally, no differentiation could be made between the different care tasks, particularly for personal and household care, partly because very few respondents received personal care. In future research that analyses more selected samples with a higher prevalence of medical problems and consequently a higher need for personal care, task differentiation will need further attention.

The findings suggest a sequence in transitions from 'no care' through 'informal care' to 'professional care'. Given the presence of a partner and a need for care, a transition to informal care seems most likely. This demands attention to the growing number of single people who will have to turn to professional care. As this may create a widening gap between the demand for professional care and its supply, the significant influence of the predisposing and enabling factors requires continued attention. Some of these factors may be amenable to interventions, not least improved communication of transparent information on care arrangements and effective individual coaching to compensate for low education. Furthermore, to achieve patterns of care delivery that are well tailored to the needs of all older people, more government financial support should be allocated to private-sector help to improve its availability.

#### Acknowledgements

This study was conducted as part of the *Longitudinal Ageing Study Amsterdam* (LASA), which is primarily funded by the Dutch Ministry of Health, Welfare and Sports and by the Netherlands Organisation for Scientific Research (NWO), Chronic Diseases Programme.

# NOTES

- <sup>1</sup> Of the 698 respondents who were lost, 18.1 per cent died before being interviewed, 56.4 per cent refused to participate, 19.2 per cent were too ill or cognitively impaired to participate, and 6.3 per cent could not be contacted. Non-response was related to age (p < 0.001) but not to sex (as was true for the non-responders at the three-yearly follow-ups).
- 2 Of the \$4\$ respondents with incomplete data, 24.8 per cent died before  $t_1$ , 49.3 per cent refused to participate, 19.3 per cent were interviewed by telephone using a much shortened instrument, 4.5 per cent were too ill or cognitively impaired to participate, and 2.1 per cent could not be contacted.
- 3 Of the 479 persons who were lost, 31.5 per cent died before t<sub>2</sub>, 49.7 per cent refused to participate, 11.1 per cent were interviewed by telephone with the shorter instrument, 5.6 per cent were too ill or cognitively impaired to participate, and 2.1 per cent could not be contacted.
- 4 In further analyses, the prevalence at  $t_0$  of cardiac disease (OR = 1.79, c.i. 1.07–2.98), peripheral arteriosclerosis (OR = 1.95, c.i. 1.04–3.63), diabetes mellitus (OR = 2.79, c.i. 1.43–5.44) and arthritis (OR = 1.72, c.i. 1.10–2.68) was associated with the transition to subsidised home-care. The transition to privately purchased home-care was associated only with the prevalence or arthritis (OR = 1.49, c.i. 1.03–2.16).
- 5 Stable functional limitations (OR = 1.92, c.i. 1.16–3.18), incident functional limitations (OR = 1.99, c.i. 1.11–3.58), incident depression (OR = 2.22, c.i. 1.15–4.28) and stable fair to poor self-rated health (OR = 1.71, c.i. 1.03–2.85) all entered as predictive of the transition to private home-care.

#### References

- Andersen, R. M. 1995. Revisiting the behavioural model and access to medical care: does it matter? *Journal of Health and Social behavior*, **35**, 1–10.
- Andersen, R. M. and Newman, J. F. 1973. Societal and individual determinants of medical care utilization in the United States. *Milbank Fund Quarterly*, **51**, 95–124.
- Berkman, L. F., Berkman, C. S., Kasl, S. V., Freeman, D. H., Leo, L., Ostield, A. M., Coroni-Huntly, J. and Brody, J. A. 1986. Depressive symptoms in relation to physical health and functioning in the elderly. *American Journal of Epidemiology*, **124**, 372–88.
- Borrayo, E. A., Salmon, J. R., Polivka, L. and Dunlop, B. D. 2002. Utilization across the continuum of long-term care services. *The Gerontologist*, **42**, 603–12.
- Broese van Groenou, M. I., van Tilburg, T. G., de Leeuw, E. D. and Liefbroer, A. C. 1995. Data collection. In Knipscheer, C. P. M., de Jong Gierveld, J., van Tilburg, T. G. and Dijkstra, P. A. (eds), *Living Arrangements and Social Networks of Older Adults, First Results.* VU University Press, Amsterdam, 185–97.
- Cantor, M. H. 1979. Neighbors and friends: an overlooked resource in the informal support system. *Research on Aging*, 1, 434–63.
- Central Bureau of Statistics 1989. *Health Interview Questionnaire*. Central Bureau of Statistics, Heerlen, The Netherlands.
- Central Bureau of Statistics 1997. Bevolkingsprognose 1996–2050 [Population projections 1996–2050]. Maandstatistiek van de Bevolking [Monthly Population Statistics], **45**, 62–70.
- Crets, S. 1996. Determinants of the use of ambulant social care by the elderly. *Social Science and Medicine*, **43**, 1709–20.
- Deeg, D. J. H. and Westendorp-de Serière, M. 1994. Autonomy and Well-Being in the Aging Population I. Report from the Longitudinal Aging Study Amsterdam 1992–1993. VU University Press, Amsterdam.

- Deeg, D. J. H., Beekman, A. T. F., Kriegsman, D. M. W. and Westendorp-de Serière, M. 1998. Autonomy and Well-Being in the Aging Population II. Report from the Longitudinal Aging Study Amsterdam 1992–1996. VU University Press, Amsterdam.
- Folstein, M. F., Folstein, S. E. and McHugh, P. R. 1975. 'Mini Mental State': a practical method for grading the cognitive state of patients for the clinician. *Journal of Psychiatric Research*, 12, 189–98.
- Geerlings, M. I., Schoevers, R. A., Beekman, A. T. F., Jonker, C., Deeg, D. J. H., Schmand, B., Ader, H. J., Bouter, L. M. and van Tilburg, W. 2000. Depression and risk of cognitive decline and Alzheimer's disease: results of two prospective communitybased studies in The Netherlands. *British Journal of Psychiatry*, **176**, 568–75.
- Geerlings, S. W., Beekman, A. T. F., Deeg, D. J. H. and van Tilburg, W. 2000. Physical health and the onset and persistence of depression in older adults: an eight-wave prospective community-based study. *Psychological Medicine*, **30**, 369–80.
- George, L. K. 1987. Easing caregiver burden: the role of informal and formal supports. In Ward, R. and Tobin, S. (eds), *Health and Aging: Sociological Issues and Policy Directions*. Springer Publishing Company, New York, 133–58.
- Glass, T. A. 1998. Conjugating the 'tenses' of function: discordance among hypothetical, experimental, and enacted function in older adults. *The Gerontologist*, **38**, 101–12.
- Hanson, E., Andersson, B. A., Magnusson, L., Lidskog, R. and Holm, K. 2002. Information Centre: responding to needs of older people and carers. *British Journal of Nursing*, 11, 935–40.
- Jacobson, N. S. and Truax, P. 1991. Clinical significance: a statistical approach to defining meaningful change in psychotherapy research. *Journal of Consulting and Clinical Psychology*, 59, 12–9.
- Jette, A. M., Tennstedt, S. L. and Branch, L. G. 1992. Stability of informal long-term care. Journal of Aging and Health, 4, 193–211.
- Kempen, G. I. and Suurmeijer, T. P. 1991. Professional home-care for the elderly: an application of the Andersen-Newman model in the Netherlands. *Social Science and Medicine*, 33, 1081–9.
- Klinkenberg M. 2004. The Last Phase of Life of Older People: Health, Preferences and Care. Unpublished Ph.D thesis, Faculty of Medicine, VU University, Amsterdam.
- Kriegsman, D. M., Penninx, B. W., van Eijk, J. T., Boeke, A. J. and Deeg, D. J. H. 1996. Self-reports and general practitioner information on the presence of chronic diseases in community dwelling elderly: a study on the accuracy of patients' self-reports and on determinants of inaccuracy. *Journal of Clinical Epidemiology*, 49, 1407–17.
- Kriegsman, D. M., van Eijk, J. T., Penninx, B. W., Deeg, D. J. H. and Boeke, A. J. 1997. Does family support buffer the impact of specific chronic diseases on mobility in community dwelling elderly? *Disability and Rehabilitation*, 19, 71–83.
- Litwak, E. 1985. Helping the Elderly: The Complementary Roles of Informal Networks and Formal ystems. Guilford, New York.
- Logan, J. R. and Spitze, G. 1994. Informal support and the use of formal services by older Americans. *Journal of Gerontology*, **49**, S25–34.
- Lyons, K. S. and Zarit, S. H. 1999. Formal and informal support: the great divide. International Journal of Geriatric Psychiatry, 14, 183–92.
- MaloneBeach, E. E., Zarit, S. H. and Spore, D. L. 1992. Caregivers' perceptions of case management and community-based services: barriers to service use. *Journal of Applied Gerontology*, 11, 146–59.
- Portrait, F. R., Lindeboom, M. and Deeg, D. J. 2000. The use of long-term care services by Dutch elderly. *Health Economics*, **9**, 513–31.
- Pot, A. M., Deeg, D. J. H. and Knipscheer, C. P. M. 2001. Institutionalization of demented elderly: the role of caregiver characteristics. *International Journal of Geriatric Psychiatry*, 16, 273–80.

- Prismant/VWS 2000. Wachten op Zorg. Wachtlijsten in de Zorg [Waiting for Care: Waiting Lists in Nursing Care]. Prismant, Utrecht, Netherlands.
- Radloff, L. S. 1977. The CES-D scale: a new self-report depression scale for research in the general population. *Applied Psychological Measurement*, **1**, 385–401.
- Roe, B. H., Whattam, M., Young, H. and Dimond, M. 2001. Elders' needs and experiences of receiving formal and informal care for their activities of daily living. *Journal of Clinical Nursing*, 10, 389–97.
- Smit, J. H., de Vries M. Z. and Poppelaars, J. L. 1998. Data collection and fieldwork procedures. In Deeg, D. J. H., Beekman, A. T. F., Kriegsman, D. M. W. and Westendorp-de Serière, M. (eds.), Autonomy and Well-Being in the Aging Population II. Report from the Longitudinal Aging Study Amsterdam 1992–1996. VU University Press, Amsterdam, 9–20.
- Sonsbeek, J. L. A. van 1988. Methodological and substantial aspects of the OECD indicator of chronic functional limitations. *Maandbericht Gezondheid*, 88, 4–17.
- Sussman, M. 1976. The family life of old people. In Binstock, R. and Shanas, E. (eds), Handbook of Aging and the Social Sciences. Van Nostrand Reinhold, New York, 415–49.
- Tennstedt, S. L., Sullivan, L. M., McKinlay, J. B. and D'Agostino, R. B. 1990. How important is functional status as a predictor of service use by older people? *Journal of Aging and Health*, 2, 439–61.
- Tennstedt, S. L., Harrow, B. S. and Sullivan, L. M. 1996. Informal care versus formal services: changing patterns of care over time. *Journal of Aging and Social Policy*, 7, 71–91.
- Tombaugh, T. N. and McIntyre, N. J. 1992. The mini-mental state examination: a comprehensive review. *Journal of the American Geriatrics Society*, **40**, 922–35.
- Ward, R. A., Sherman, S. and LaGory, M. 1984. Informal networks and knowledge of services for older persons. *Journal of Gerontology*, 39, 216–23.
- Zarit, S. H., Gaugler, J. E. and Jarrott, S. E. 1999. Useful services for families: research findings and directions. *International Journal of Geriatric Psychiatry*, 14, 165–77.

Accepted 7 June 2004

Address for correspondence:

Sandra W. Geerlings, Longitudinal Ageing Study Amsterdam, Van der Boechorststraat 7, 1081 BT Amsterdam, The Netherlands.

e-mail: sw.geerlings@vumc.nl