Title

Association between Dementia, Change in Home-care Use, and Depressive Symptoms during the COVID-19 Pandemic: A Longitudinal Study Using Data from Three Cohort Studies

Authors

Miharu Nakanishi, RN, PhD^{a,b,c*}, Syudo Yamasaki, CPT, PhD^c, Taeko Nakashima, SW, PhD^d, Yuki Miyamoto, RN, PhD^e, Claudia Cooper, MD, PhD^f, Marcus Richards, MD, PhD^g, Daniel Stanyon, MSc^c, Mai Sakai, RN, PhD^b, Hatsumi Yoshii, RN, PhD^b, and Atsushi Nishida, PSW, PhD^c

Affiliations

^a Department of Public Health and Primary Care, Leiden University Medical Center, Hippocratespad 21, 2333 ZD Leiden, the Netherlands

^b Department of Psychiatric Nursing, Tohoku University Graduate School of Medicine, 2-1 Seiryo-machi, Aoba-ku, Sendai-shi, Miyagi, 980-8575 Japan

^c Research Center for Social Science & Medicine, Tokyo Metropolitan Institute of Medical

Science, 2-1-6 Kamikitazawa, Setagaya-ku, Tokyo, 156-8506 Japan

^d Department of Social Healthcare and Business, Faculty of Healthcare Management, Nihon

Fukushi University, Mihama-cho, Aichi, 470-3295 Japan

^e Department of Psychiatric Nursing, School of Health Sciences and Nursing, Graduate
 School of Medicine, The University of Tokyo, Bunkyo-ku, Tokyo, 113-0033 Japan
 ^f Centre for Psychiatry and Mental Health, Wolfson Institute of Population Health, Queen
 Mary University of London, London, E1 2AB the United Kingdom
 ^g MRC Unit for Lifelong Health & Ageing at UCL, University College London, London,
 WC1E 7HB the United Kingdom

Running title

Home care and depression in dementia

*Correspondence to:

Miharu Nakanishi PhD, Department of Public Health and Primary Care, Leiden University

Medical Center, Hippocratespad 21, 2333 ZD Leiden, the Netherlands.

Telephone: +31 71 526 84 81

Email: mnakanishi-tky@umin.ac.jp

Abstract (248/250)

Background: The emotional impact of the coronavirus disease 2019 (COVID-19) pandemic

on people with dementia has been quantified. However, little is known about the impact of change in home-care use owing to the pandemic.

Objective: To determine the longitudinal association between dementia, change in home-care use, and depressive symptoms during the pandemic.

Methods: We included data of 43,782 home-dwelling older adults from the English Longitudinal Study of Ageing (ELSA), Study of health, Ageing and Retirement in Europe (SHARE), and National Health and Aging Trends Study (NHATS). This study considered the latest main wave survey prior to the pandemic as the baseline, and the COVID-19 survey as follow-up. In a series of coordinated analyses, multilevel binomial logistic regression model was used to examine the association between baseline dementia, change in home-care use at follow-up, and presence of depressive symptoms.

Results: Dementia, using the ELSA, SHARE, and NHATS datasets, was identified in 2.9%, 2.3%, and 6.5% of older adults, and home-care use reduced in 1.7%, 2.8%, and 1.1% of individuals with dementia, respectively. Dementia was significantly associated with the increased risk of depressive symptoms in all three cohorts. However, the interaction between dementia and period (follow-up) was non-significant in SHARE and NHATS. Across all three cohorts, home-care use during the pandemic, regardless of change in amount, was significantly associated with increased depressive symptoms, compared to the non-use of home care.

Conclusion: These results highlight the need for tailoring dementia care at home to promote independence and provide sustainable emotional support.

Keywords: cohort studies; dementia; depression; home care services; SARS-CoV 2; social

interaction

INTRODUCTION

2	The coronavirus disease pandemic, which began in 2019 (COVID-19) was caused by
3	severe acute respiratory syndrome coronavirus 2 (SARS-CoV 2) [1]. It particularly affected
4	older populations, especially persons with dementia. The morbidity and mortality of COVID-
5	19 worsen with advancing age and pre-existing health conditions; therefore, many persons
6	with dementia face a high risk [2]. Moreover, persons with dementia may be especially
7	susceptible to the social consequences of lockdown and confinement [3], including loss of
8	support from primary caregivers and restricted social interactions [4-6]. Furthermore, persons
9	with dementia are more likely to rely on formal care services for social contact as well as
10	practical support [7]. Even before the pandemic, persons with dementia and caregivers faced
11	challenges with respect to social interactions and mental health issues [8]. Once restrictions
12	were imposed during the COVID-19 pandemic, social contact was greatly reduced, which
13	caused negative psychological and cognitive effects [9-11]. Several studies have quantified
14	emotional changes such as depression and anxiety from before the pandemic to during the
15	pandemic among people with dementia [12–15]. Furthermore, recent studies based on data
16	from population-based cohorts in England [16] and the United States [17] have shown that
17	persons with dementia experienced worse mental health outcomes during the pandemic
18	compared to those without the condition. However, these studies did not measure the impact
19	of change in home-care use due to the COVID-19 pandemic.

20	We aimed to investigate the relationship between dementia, change in home-care use,
21	and change in depressive symptoms before and during the COVID-19 pandemic. We used
22	three population-based cohorts to include older adults with probable dementia which may not
23	necessarily be identified through clinical diagnoses. We hypothesized that compared with
24	older adults without impairment, those with dementia would show a greater increase in
25	depressive symptoms during the pandemic. Additionally, we hypothesized that any increases
26	in depressive symptoms during the pandemic would be reduced in magnitude by controlling
27	for home care, since home-care workers have the potential to provide social and practical
28	support.
29	
30	MATERIALS AND METHODS
31	Study Design and Setting
31 32	Study Design and Setting We used data obtained from three longitudinal studies of aging: the English Longitudinal
31 32 33	Study Design and Setting We used data obtained from three longitudinal studies of aging: the English Longitudinal Study of Ageing (ELSA) [18], the Study of Health, Ageing and Retirement in Europe
31 32 33 34	 Study Design and Setting We used data obtained from three longitudinal studies of aging: the English Longitudinal Study of Ageing (ELSA) [18], the Study of Health, Ageing and Retirement in Europe (SHARE) [19], and the National Health and Aging Trends Study (NHATS) [20].
3132333435	Study Design and Setting We used data obtained from three longitudinal studies of aging: the English Longitudinal Study of Ageing (ELSA) [18], the Study of Health, Ageing and Retirement in Europe (SHARE) [19], and the National Health and Aging Trends Study (NHATS) [20]. ELSA is a nationally representative sample of men and women aged ≥50 years who
 31 32 33 34 35 36 	Study Design and Setting We used data obtained from three longitudinal studies of aging: the English Longitudinal Study of Ageing (ELSA) [18], the Study of Health, Ageing and Retirement in Europe (SHARE) [19], and the National Health and Aging Trends Study (NHATS) [20]. ELSA is a nationally representative sample of men and women aged ≥50 years who were resident in England [18]. It began in 2002 (wave 1), and the assessment is repeated
 31 32 33 34 35 36 37 	Study Design and Setting We used data obtained from three longitudinal studies of aging: the English Longitudinal Study of Ageing (ELSA) [18], the Study of Health, Ageing and Retirement in Europe (SHARE) [19], and the National Health and Aging Trends Study (NHATS) [20]. ELSA is a nationally representative sample of men and women aged ≥50 years who were resident in England [18]. It began in 2002 (wave 1), and the assessment is repeated every 2 years. The current study drew the pre-pandemic responses from the main ELSA wave

39	of ELSA, and performed in two waves (June/July and November/December 2020). During
40	the pandemic, COVID-related restrictions meant that ELSA's usual face-to-face interview
41	approach could not implemented. Instead, the COVID-19 waves used a sequential mixed-
42	mode design with an online survey and telephonic interviewing. During the first COVID-19
43	survey, due to a survey error, about 75% of respondents were not asked the eighth depression
44	item [16]. Therefore, we used data from the second COVID-19 survey for the assessment of
45	the pandemic period. The South Central – Berkshire Research Ethics Committee provided
46	ethical approval for ELSA (21/SC/0030, March 22, 2021).
47	SHARE included data on the lives of Europeans aged 50+ years from 28 countries
48	including Israel [19]. SHARE is closely modelled after and constantly harmonized with its
49	sister studies including ELSA. Usually, data are collected biannually based on computer-
50	assisted personal interviewing (CAPI). The interviewers conduct face-to-face interviews
51	using a laptop on which the CAPI instrument is installed. Owing to the COVID-19 outbreak,
52	data collection and the fieldwork were suspended in March 2020 after 70% of the regular
53	interviews of SHARE wave 8 survey (October 2019-March 2020) were performed, which
54	was conducted in 27 countries (excluding Portugal) [21,22]. A switch to telephone
55	administered interviews (CATI) was decided as the alternative to the previous face-to-face
56	interviewing, and the first SHARE Corona Survey was conducted from June to September
57	2020 [23,24]. The Ethics Committee of the University of Mannheim and the Ethics Council

58 of the Max Planck Society provided ethical approval for SHARE.

59	NHATS is a nationally representative sample of adults aged ≥ 65 who are Medicare
60	beneficiaries in the United States [20]. Although there is another cohort for aging, namely the
61	Health and Retirement Study (HRS), a sister study of ELSA and SHARE, the HRS data
62	collection during the pandemic was performed as the main wave survey between March 2020
63	and May 2021. Considering the comparability across cohorts, we chose NHATS from the
64	cohort studies in the United States. Data collection of NHATS began in 2011, and the
65	replacement of the study sample occurred in 2015. Annual interviews are administered to
66	members of NHATS. At the onset of the COVID-19 pandemic, research that relied on in-
67	person contact was widely prohibited. Consequently, the interview was conducted by
68	telephone in Round 10. The COVID-19 questionnaires were mailed from June 2020 ending
69	through October 2020 ending, following the round-10 collection. Pre-pandemic responses
70	were drawn from the 2019 NHATS round 9 survey. The Johns Hopkins Bloomberg School of
71	Public Health IRB provided ethical approval for NHATS.
72	For each cohort study, we considered the main survey prior to the pandemic as the
73	baseline, and COVID-19 survey as the follow-up. Procedures involving experiments on
74	human subjects are done in accord with the Helsinki Declaration of 1975.
75	

77	We selected participants from the three cohorts if they (1) were adults aged 50 years or
78	older at baseline, (2) were living in a private home at baseline, (3) had participated in the
79	COVID-19 survey, and (4) were home-dwelling adults during the pandemic (Figure 1).
80	
81	Measurements
82	We conducted a coordinated, identical analyses across the different datasets to examine
83	relationships across samples using conceptually equivalent measures of the constructs of
84	interest [25]. Although measures differed across studies, the same covariates, measurement
85	scores, and modeling could be used to obtain comparable results across datasets, and identify
86	sources of discrepancy across studies.
87	Multiple variables were recoded to create equivalent versions across all datasets, as
88	described below (see Supplementary Table S1). This allows a comparison of results across
89	models to inform substantive conclusions.
90	
91	Dementia
92	Considering the prevalent clinical misdiagnosis of dementia [26], we used the presence
93	of dementia as the primary independent variable, irrespective of any clinical dementia
94	diagnosis. We used the term "dementia" for probable dementia case throughout. Dementia
95	caseness was determined by algorithms assessing the likelihood of a dementia diagnosis,

96 rather than relying on clinical diagnosis.

97	For ELSA and SHARE, we applied a common classification approach for dementia,
98	consistent with previous studies on dementia in SHARE [27-29]. In each wave, participants
99	were asked to complete a memory recall task (immediate and delayed recall of 10 common
100	words) and an animal fluency task (naming as many animals as possible in 60 seconds) [30].
101	Participants who performed 1.5 standard deviations (SD) below the age-graded (5-year) mean
102	of immediate or delayed recall (or both) were coded as 1 and compared to other participants,
103	coded 0. Similarly, those who performed 1.5 SD below the age-graded mean of verbal
104	fluency were coded as 1, compared to others coded 0. Dementia was defined as a score of 1
105	on both the memory and verbal fluency tasks.
106	In the NHATS, dementia was identified using a validated algorithm for surveys [31]
107	based on a report of either dementia or Alzheimer's disease (diagnosed by a physician),
108	cognitive tests of memory, orientation, and executive function. These were used to form a
109	derived variable that was calculated using the eight-item responses to Differentiate Aging and
110	Dementia (AD8) instrument [32]. We used the algorithm because an animal fluency task was
111	not employed in the NHATS and we could not apply the classification approach for ELSA
112	and SHARE as mentioned above.

Depressive Symptoms

115	We defined depressive symptoms as the mental health outcomes assessed at baseline and
116	follow-up. Since the assessment in SHARE was based on binary response options (yes or no),
117	we used the presence of clinically significant symptoms as the independent variable for the
118	main analyses across the three cohorts.
119	Depressive symptoms in ELSA were measured using the shortened version of the Center
120	for Epidemiologic Studies Depression (CESD) scale [33]. The threshold for clinically
121	significant symptoms was four or more positive symptoms.
122	In SHARE main wave surveys, depressive symptoms were assessed using a self-
123	reported European-Depression (EURO-D) scale [34]. However, SHARE Corona Survey
124	adapted only two items (depression and sleep) from EURO-D. In this study, we used a single
125	question for depression with a yes/no response option: "In the last month, have you been sad
126	or depressed?"
127	Depression in NHATS was assessed using the Patient Health Questionnaire-2 (PHQ-2)
128	[35]. The PHQ-2, a validated two-question-screening questionnaire, clarifies the frequency of
129	depressed mood and anhedonia during the preceding two weeks. The total score ranges from
130	0 to 6 while the threshold for substantial depressive symptoms was ≥ 3 [36].
131	
132	Changes in Home-care Use During the COVID-19 Pandemic

133 During the COVID-19 pandemic, changes in home-care use was typically assessed using

134	multiple questions about (1) whether participants recently received home care, and if yes, (2)
135	whether amount of care was less, same, or greater than it was before the pandemic. In this
136	study, we reclassified participants into three categories: (i) older adults who did not receive
137	home care (no home care used), (ii) those who received same or increased amount of care
138	compared with before the pandemic (same or increased), and (iii) those who received reduced
139	amount of care than before. In SHARE, participants were asked whether they faced more
140	difficulties in receiving the amount of home care that they needed. If they answered "yes,"
141	they were also asked whether they had to pay more to receive the help they needed, or
142	whether those who cared for them could not come to their home. Therefore, we assumed that
143	participants who answered "yes" to the former question may have experienced reduced
144	amount of care during the pandemic.
145	
146	Sociodemographic characteristics
147	We considered the following sociodemographic characteristics: age, sex, whether the
148	respondents had a spouse or partner, living alone, and educational attainment. The
149	aforementioned variables were selected as potentially relevant to mental health outcomes
150	among older adults during the pandemic across the three cohorts [17, 37-40]. Given the
151	diversity in measurements and definitions across the three cohorts, we opted to exclude
152	socioeconomic status, alcohol consumption, and physical comorbidities in this study,

although these variables could have been relevant to depressive symptoms.

154

155 Statistical Analysis

156	First, we calculated the descriptive statistics across the three cohorts. Thereafter, we
157	performed multilevel binomial logistic regression analyses of the presence of depressive
158	symptoms. A panel-data format was adopted wherein the same older adult appeared two
159	times. The first model used presence of depressive symptoms as the dependent variable, and
160	period (baseline or follow-up), dementia, change in home-care use, and interaction between
161	dementia and period as the independent variables. All sociodemographic variables were also
162	included as independent variables. Change in home-care use, period, and interaction term
163	were treated as within-person time-variant variables, and dementia and sociodemographic
164	variables were entered as between-person time-invariant variables. A sensitivity analysis of
165	the first model was conducted for ELSA and NHATS using a four-category classification of
166	home-care use (no home care used, reduced, same, or increased). Since the prevalence of
167	reduced home-care use was low (0.5–2.8%), the first model was reanalyzed excluding
168	participants with a reduced amount of care from the sample across the three cohorts.
169	The second model added interaction terms between dementia and change in home-care
170	use as the within-person variables to the first model. Given the potential bias in logistic
171	regression models [41, 42] and the missing evaluation for increased depressive symptoms

172	under the threshold, another sensitivity analysis of the second model was performed for
173	ELSA and NHATS using linear regression analyses of total scores of depressive symptoms as
174	the independent variable.
175	In the regression analysis, we used the full information maximum likelihood to handle
176	missing data [43]. Data management was conducted using Stata 18.0 (StataCorp). We
177	performed regression analyses using Mplus for Windows, version 8.10 (Muthén & Muthén,
178	Los Angeles, CA, USA). Statistical significance was set at $\alpha = 0.05$.
179	
180	RESULTS
181	A total of 43,782 older adults aged \geq 50 years were included in this study. Table 1
182	displays baseline characteristics of participants per cohort. NHATS included more older
183	adults aged ≥85 years (23.4%) than ELSA (2.9%) or SHARE (7.4%). Overall, 1,146
184	individuals were identified with probable dementia at baseline (2.7%) including 2.9% in
185	ELSA (N = 6,114), 2.3% in SHARE (N = 33,263), and 6.5% in NHATS (N = 3,001) (Table
186	1).
187	Table 2 shows the numbers and percentages of changes in amount of home care per
188	cohort. During the pandemic, more than 90% of participants in ELSA and SHARE did not
189	receive home care. Approximately 16–17% of individuals with dementia received home care
190	in the two cohorts. Contrarily, 1,398 (48.9%) participants in NHATS received home care

191	during the pandemic, including 79% of individuals with dementia received home care. Few
192	participants (0.5–2.8%) reported reduction in amount of care since the COVID-19 outbreak,
193	regardless of dementia status (Table 2).
194	Figure 2 indicates the presence of depressive symptoms at baseline and follow-up by
195	dementia status per cohort. Across all three cohorts, individuals with dementia appeared to
196	have higher percentages of depressive symptoms than those without impairment. In ELSA
197	and NHATS, depressive symptoms increased during the pandemic than before. However, the
198	decline in depressive symptoms was more observed in SHARE participants, as is described in
199	a previous study [44] (Figure 2, Supplementary Table S2).
200	Table 3 and Supplementary Table S3 summarize results of the first multilevel binomial
201	logistic regression analyses models. Across all three cohorts, dementia was significantly
202	associated with increased risk of depressive symptoms. In ELSA, participants with dementia
203	were less likely to report presence of depressive symptoms during the pandemic. However,
204	the interaction between dementia and period was non-significant in SHARE and NHATS.
205	Regardless of whether there was a reduction, or same or increase in care amount, participants
206	who received home care were more likely to report presence of depressive symptoms than
207	those with no home-care use (Table 3, Supplementary Table S3). A sensitivity analysis using
208	the four-category variable of home-care use revealed that a reduced or the same amount of
209	home-care use was significantly associated with an increased risk of depressive symptoms.

210	While increased home-care use was significantly associated with an increased risk of
211	depressive symptoms in NHATS, this association was not observed in ELSA (Supplementary
212	Table S4). Another sensitivity analysis excluding participants with reduced home-care use did
213	not alter the association between increased or the same amount of home-care use and
214	depressive symptoms in all the three cohorts (Supplementary Table S5).
215	Table 4 and Supplementary Table S6 present the results of the second model wherein
216	interaction terms were added to the first model. In ELSA, participants with dementia and
217	home-care use regardless of change in amount were more likely to report presence of
218	depressive symptoms. However, in SHARE and NHATS, interactions between dementia and
219	change in home-care use were non-significant (Table 4, Supplementary Table S6). A
220	sensitivity analysis using total scores of depressive symptoms did not significantly change the
221	results (Supplementary Table S7).
222	
223	Discussion
224	To our knowledge, this is the first study on a general population that examined
225	association between dementia, change in home-care use, and depressive symptoms during the
226	pandemic. Numerous studies have demonstrated worsening mental health in persons with
227	dementia [9–15]. A few studies further revealed that those with dementia were at higher risk
228	of depressive symptoms than the general older adults during the pandemic [16,17]. However,

229	the association between dementia and change in depressive symptoms was inconsistent
230	across the three cohorts. The interaction effects between dementia and period and that
231	between dementia and home-care reduction were observed in ELSA, but not in either
232	SHARE or NHATS. Contrary to our hypothesis, receiving home care was consistently
233	associated with worse depressive symptoms across the three cohorts. This is the first study to
234	demonstrate a consistent association between home-care use during the pandemic and worse
235	depressive symptoms across England, other 27 European countries, and the United States.
236	Older adults who did not use home care were less likely to report depressive symptoms
237	during the pandemic than those who used home care, regardless of dementia status or change
238	in care amount.
239	Participants with dementia were more likely to receive care at home during the
240	pandemic. Receiving care at home was related to worse depressive symptoms, which is
241	consistent with a previous report that activity of daily living (ADL) and mobility impairment
242	were associated with worse mental health changes [38]. These findings suggest that the loss
243	of independence among older adults may have exacerbated the impact of pandemic-related
244	restrictions and reduced social interactions on mental health. As persons with dementia are at
245	increased risk for COVID-19 infection [2], they were particularly targeted for compliance
246	with physical distancing measures, which likely result in reduced social interactions [45-47].
247	Considering that friendships, sense of belonging, and feeling valued within social

248	connections are important for mental health in older adults [46], individuals with less
249	resilience for substituting their loss of social contacts may have magnified adverse effects
250	[47]. Our findings indicate that in any similar future pandemic, emotional support for social
251	connectedness should be integrated and sustained for home-care users with dementia
252	wherever possible, and that potentially significant detriments to mental health need to be
253	weighed against risks of infection and physical illness [48–51]. In the early pandemic
254	months, numerous individuals transitioned from in-person to video contact in order to
255	maintain social ties [52]. This type of transition may have been challenging among older
256	adults with dementia. Despite campaigns by the Alzheimer's Society United Kingdom and
257	other charities [53], home-care agencies that were not specialized in dementia care, as well as
258	family and friends who were non-caregivers, compared with caregivers, may have been less
259	aware of these campaigns. Therefore, additional strategies should be explored to achieve this
260	goal.
261	In our study, changes in amount of care due to the pandemic substantially varied across
262	England, other 27 European countries, and the United States. In the United States, most older
263	adults with dementia received same or increased amount of home care. In England and other
264	27 European countries, more than 80% of participants with dementia did not receive home
265	care during the pandemic. As participants in ELSA and SHARE were younger than those in
266	NHATS, the cross-cohort difference in home-care use could have reflected that these younger

267	participants had fewer care needs, resulting in refraining from home-care use during the
268	pandemic. However, we did not consider baseline home-care use before the pandemic, as the
269	questions and definitions of home care were typically different from those in the COVID-19
270	study in each cohort. Some participants classified into "no home-care use" could have
271	belonged to the "reduced" cases who had been home-care users before the pandemic.
272	Furthermore, it should be noted that data about home-care use in SHARE and NHATS could
273	not be used to calculate the proportion of informal and formal care. It is possible that the
274	family and friends of older adults with dementia might have compensated for the decreased
275	formal care, which has been suggested by studies during the early pandemic months [54,55].
276	This shift in care could have resulted in reduced social networks and contacts [56], which
277	could have increased the adverse effects of the pandemic on mental health outcomes in
278	individuals with dementia.
279	
280	Strengths and Limitations
281	The main strength of this study is the use of representative cohorts from England, other
282	27 European countries, and the United States. The longitudinal design allowed for
283	comparisons of outcome measures before and during the pandemic. Further, our study

- 284 included a sufficient number of individuals with dementia for statistical comparison with
- those without impairment. The use of full information maximum likelihood estimation

286	enabled us to include participants with missing data in the multilevel binomial logistic
287	regression analyses. The findings obtained across the three cohorts provided insights into the
288	consistent experiences and needs of individuals with dementia after long-term restrictions,
289	regardless of country-specific healthcare systems and COVID-19 related counter-measures.
290	However, each cohort used different depression metrics such that the direct comparisons
291	of depressive symptoms across cohorts were limited. The degree of comparability would also
292	be compromised by the cross-cohort differences in the prevalence of dementia and change in
293	home-care use. Participants in ELSA and SHARE were younger than those in NHATS; the
294	difference in age distribution may have led to relatively small number of participants with
295	dementia in those cohorts. Similarly, participants in NHATS experienced care reduction
296	during the pandemic less frequently. Cognitive function during the pandemic might have
297	been subject to the risk of decline, which could have led to a reverse causation between
298	dementia and depressive symptoms. In addition, measures regarding resilience that may have
299	helped people with dementia adapt to challenges [57,58] were not included in this study.
300	Notably, we did not consider social contact with family or friends, which may prevent mental
301	health deterioration. Although COVID-19 sub-studies usually assessed social contact with
302	family and friends, it often involved different measures from those in main wave surveys.
303	Additionally, social contact with family or friends may overlap with informal care at home
304	during the pandemic. Individuals with severe dementia may have difficulties answering

305	surveys, though the measures (e.g., trained interviewers) were taken to mitigate it. We did not
306	have data on prescriptions or dysphasia, which may affect performance in cognitive tests.
307	Furthermore, some risk factors for depressive symptoms, such as socioeconomic status,
308	alcohol consumption, or physical comorbidities were not included in this study due to the
309	diversity in measurement across the three cohorts. Finally, the participants in this study only
310	had two times of assessment. Future follow-up research to evaluate mental health changes
311	would be helpful to project the trajectory.
312	
313	
0 - 0	Conclusions
314	Conclusions Our findings demonstrate that older adults who received home care, regardless of
314 315	Conclusions Our findings demonstrate that older adults who received home care, regardless of change in care amount and dementia status, during the outbreak were more likely to report
314 315 316	Conclusions Our findings demonstrate that older adults who received home care, regardless of change in care amount and dementia status, during the outbreak were more likely to report worse depressive symptoms compared with those who received no care at home. The loss of
314315316317	Conclusions Our findings demonstrate that older adults who received home care, regardless of change in care amount and dementia status, during the outbreak were more likely to report worse depressive symptoms compared with those who received no care at home. The loss of independence among older adults may have exacerbated the impact of pandemic-related
 314 315 316 317 318 	Conclusions Our findings demonstrate that older adults who received home care, regardless of change in care amount and dementia status, during the outbreak were more likely to report worse depressive symptoms compared with those who received no care at home. The loss of independence among older adults may have exacerbated the impact of pandemic-related related restrictions and reduced social interactions on mental health. As people with dementia
 314 315 316 317 318 319 	Conclusions Our findings demonstrate that older adults who received home care, regardless of change in care amount and dementia status, during the outbreak were more likely to report worse depressive symptoms compared with those who received no care at home. The loss of independence among older adults may have exacerbated the impact of pandemic-related related restrictions and reduced social interactions on mental health. As people with dementia are more likely to receive care at home, there is a need for promoting independence and

321 ACKNOWLEDGEMENTS

322 The authors have no acknowledgments to report.

323

324 FUNDING

325	This work was supported by the Ministry of Education, Culture, Sports, Science, and
326	Technology of Japan under Grant [number JP21H05173]; Japan Society for the Promotion of
327	Science [numbers JP21H03281; JP22KK0258]; Japan Agency for Medical Research and
328	Development under Grant [number 22579506]; and the UK Medical Research Council under
329	Grant [numbers MC_UU_10019/1 and /3]. This work was partly supported by the Tohoku
330	University Graduate School of Medicine. They had no input into data collection,
331	management, analysis, or interpretation, and were not able to monitor the manuscript for
332	presentation, review, or approval.
333	The English Longitudinal Study of Ageing (ELSA) was supported by the National
334	Institute on Aging (Grant R01AG017644) and by a consortium of UK government
335	departments coordinated by the National Institute for Health Research. Funding for data
336	collection during the COVID-19 pandemic was provided by the UK Research and Innovation
337	Economic and Social Research Council COVID-19 Rapid Response Initiative
338	(ES/V003941/1).

339 The Study of Health, Ageing and Retirement in Europe (SHARE) data collection has

- been funded by the European Commission, DG RTD through FP5 (QLK6-CT-2001-00360),
- 341 FP6 (SHARE-I3: RII-CT-2006-062193, COMPARE: CIT5-CT-2005-028857, SHARELIFE:
- 342 CIT4-CT-2006-028812), FP7 (SHARE-PREP: GA N°211909, SHARE-LEAP: GA
- 343 N°227822, SHARE M4: GA N°261982, DASISH: GA N°283646) and Horizon 2020
- 344 (SHARE-DEV3: GA N°676536, SHARE-COHESION: GA N°870628, SERISS: GA
- 345 N°654221, SSHOC: GA N°823782, SHARE-COVID19: GA N°101015924) and by DG
- 346 Employment, Social Affairs & Inclusion through VS 2015/0195, VS 2016/0135, VS
- 347 2018/0285, VS 2019/0332, and VS 2020/0313. Additional funding from the German Ministry
- of Education and Research, the Max Planck Society for the Advancement of Science, the U.S.
- 349 National Institute on Aging (U01_AG09740-13S2, P01_AG005842, P01_AG08291,
- 350 P30_AG12815, R21_AG025169, Y1-AG-4553-01, IAG_BSR06-11, OGHA_04-064,
- 351 HHSN271201300071C, RAG052527A) and from various national funding sources is
- 352 gratefully acknowledged (see www.share-project.org).
- 353 The National Health and Ageing Trend Study (NHATS) was produced and distributed by
- 354 www.nhats.org with funding from the National Institute on Aging (grant number
- 355 U01AG032947).

356

357 CONFLICT OF INTEREST

358 The authors have no conflict of interest to report. MN is an Editorial Board Member of

this journal, but was not involved in the peer-review process nor had access to any

360 information regarding its peer-review.

361

362 AUTHOR CONTRIBUTIONS

- 363 M.N. performed conceptualization, data curation, formal analysis, funding acquisition,
- investigation, methodology, visualization, and writing original draft. S.Y., T.N., and Y.M.
- 365 contributed to formal analysis, funding acquisition, methodology, validation, and writing –
- 366 review & editing. C.C., M.R., and D.S. contributed to conceptualization, data curation, and
- investigation, and provided resources, supervision, and writing review & editing. S.M.,
- 368 Y.H., and A.N. conducted project administration, and provided software, supervision, and
- 369 writing review & editing.

370

371 DATA AVAILABILITY STATEMENT

- 372 The data used in this study were available from the following sources:
- 373 English Longitudinal Study of Ageing (ELSA): the UK Data Service with access codes SN
- 8688, and 5050; Study of Health, Ageing and Retirement in Europe (SHARE):
- 375 http://www.share-project.org; and National Health and Aging Trends Study (NHATS):
- 376 https://www.nhats.org/researcher.

377 REFERENCES

378	[1]	Kola L, Kohrt BA, Hanlon C, Naslund JA, Sikander S, Balaji M, Benjet C, Cheung
379		EYL, Eaton J, Gonsalves P, Hailemariam M, Luitel NP, Machado DB, Misganaw E,
380		Omigbodun O, Roberts T, Salisbury TT, Shidhaye R, Sunkel C, Ugo V, van Rensburg
381		AJ, Gureje O, Pathare S, Saxena S, Thornicroft G, Patel V (2021) COVID-19 mental
382		health impact and responses in low-income and middle-income countries: reimagining
383		global mental health. Lancet Psychiatry 8, 535–550.
384	[2]	Livingston G, Huntley J, Sommerlad A, Ames D, Ballard C, Banerjee S, Brayne C,
385		Burns A, Cohen-Mansfield J, Cooper C, Costafreda SG, Dias A, Fox N, Gitlin LN,
386		Howard R, Kales HC, Kivimäki M, Larson EB, Ogunniyi A, Orgeta V, Ritchie K,
387		Rockwood K, Sampson EL, Samus Q, Schneider LS, Selbæk G, Teri L, Mukadam N
388		(2020) Dementia prevention, intervention, and care: 2020 report of the Lancet
389		Commission. Lancet 396 , 413–446.
390	[3]	Suárez-González A, Rajagopalan J, Livingston G, Alladi S (2021) The effect of
391		COVID-19 isolation measures on the cognition and mental health of people living with
392		dementia: A rapid systematic review of one year of quantitative evidence.
393		EClinicalMedicine 39 , 101047.
394	[4]	Giebel C, Cannon J, Hanna K, Butchard S, Eley R, Gaughan A, Komuravelli A,

395 Shenton J, Callaghan S, Tetlow H, Limbert S, Whittington R, Rogers C, Rajagopal M,

396		Ward K, Shaw L, Corcoran R, Bennett K, Gabbay M (2021) Impact of COVID-19
397		related social support service closures on people with dementia and unpaid carers: a
398		qualitative study. Aging Ment Health 25, 1281–1288.
399	[5]	Giebel C, Lord K, Cooper C, Shenton J, Cannon J, Pulford D, Shaw L, Gaughan A,
400		Tetlow H, Butchard S, Limbert S, Callaghan S, Whittington R, Rogers C, Komuravelli
401		A, Rajagopal M, Eley R, Watkins C, Downs M, Reilly S, Ward K, Corcoran R, Bennett
402		K, Gabbay M (2021) A UK survey of COVID-19 related social support closures and
403		their effects on older people, people with dementia, and carers. Int J Geriatr Psychiatry
404		36 , 393–402.
405	[6]	Giebel C, Pulford D, Cooper C, Lord K, Shenton J, Cannon J, Shaw L, Tetlow H,
406		Limbert S, Callaghan S, Whittington R, Rogers C, Komuravelli A, Rajagopal M, Eley
407		R, Downs M, Reilly S, Ward K, Gaughan A, Butchard S, Beresford J, Watkins C,
408		Bennett K, Gabbay M (2021) COVID-19-related social support service closures and
409		mental well-being in older adults and those affected by dementia: a UK longitudinal
410		survey. BMJ Open 11, e045889.
411	[7]	Daley S, Akarsu N, Armsby E, Farina N, Feeney Y, Fine B, Hughes L, Pooley J, Tabet
412		N, Towson G, Banerjee S (2022) What factors have influenced quality of life in people
413		with dementia and their family carers during the COVID-19 pandemic: a qualitative
414		study. BMJ Open 12, e053563.

415	[8]	Balouch S, Rifaat E, Chen HL, Tabet N (2019) Social networks and loneliness in
416		people with Alzheimer's dementia. Int J Geriatr Psychiatry 34, 666–673.
417	[9]	Liu KY, Howard R, Banerjee S, Comas-Herrera A, Goddard J, Knapp M, Livingston G,
418		Manthorpe J, O'Brien JT, Paterson RW, Robinson L, Rossor M, Rowe JB, Sharp DJ,
419		Sommerlad A, Suárez-González A, Burns A (2021) Dementia wellbeing and COVID-
420		19: Review and expert consensus on current research and knowledge gaps. Int J Geriatr
421		Psychiatry 36, 1597–1639.
422	[10]	Tsapanou A, Papatriantafyllou JD, Yiannopoulou K, Sali D, Kalligerou F, Ntanasi E,
423		Zoi P, Margioti E, Kamtsadeli V, Hatzopoulou M, Koustimpi M, Zagka A,
424		Papageorgiou SG, Sakka P (2021) The impact of COVID-19 pandemic on people with
425		mild cognitive impairment/dementia and on their caregivers. Int J Geriatr Psychiatry
426		36 , 583–587.
427	[11]	Tuijt R, Frost R, Wilcock J, Robinson L, Manthorpe J, Rait G, Walters K (2021) Life
428		under lockdown and social restrictions - the experiences of people living with dementia
429		and their carers during the COVID-19 pandemic in England. BMC Geriatr 21, 301.
430	[12]	Tsugawa A, Sakurai S, Inagawa Y, Hirose D, Kaneko Y, Ogawa Y, Serisawa S,
431		Takenoshita N, Sakurai H, Kanetaka H, Hirao K, Shimizu S (2020) Awareness of the
432		COVID-19 outbreak and resultant depressive tendencies in patients with severe
433		Alzheimer's disease. J Alzheimers Dis 77, 539–541.

434	[13]	Borelli WV, Augustin MC, de Oliveira PBF, Reggiani LC, Bandeira-de-Mello RG,
435		Schumacher-Schuh AF, Chaves MLF, Castilhos RM (2021) Neuropsychiatric
436		symptoms in patients with dementia associated with increased psychological distress in
437		caregivers during the COVID-19 pandemic. J Alzheimers Dis 80, 1705–1712.
438	[14]	Pongan E, Dorey JM, Borg C, Getenet JC, Bachelet R, Lourioux C, Laurent B,
439		COVCARE Group, Rey R, Rouch I (2021) COVID-19: association between increase of
440		behavioral and psychological symptoms of dementia during lockdown and caregivers'
441		poor mental health. J Alzheimers Dis 80, 1713–1721.
442	[15]	Tsiakiri A, Vlotinou P, Terzoudi A, Heliopoulos I, Vadikolias K (2022) Cognitive,
443		functional, and emotional changes during the COVID-19 pandemic in Greek patients
444		with neurocognitive disorders. J Alzheimers Dis 88, 537-547.
445	[16]	Beach B, Steptoe A, Zaninotto P (2023) Depression and anxiety in people with
446		cognitive impairment and dementia during the COVID-19 pandemic: Analysis of the
447		English Longitudinal Study of Ageing. PLoS Med 20, e1004162.
448	[17]	Nakanishi M, Ogawa A, Sakai M, Yoshii H, Miyashita M, Yamasaki S, Nishida A
449		(2023) Depression and anxiety in older adults with dementia during the COVID-19
450		pandemic. J Alzheimers Dis Rep 7, 307–315.
451	[18]	Steptoe A, Breeze E, Banks J, Nazroo J (2013) Cohort profile: the English Longitudinal
452		Study of Ageing. Int J Epidemiol 42, 1640–1648.

453	[19]	Börsch-Supan A, Brandt M, Hunkler C, Kneip T, Korbmacher J, Malter F, Schaan B,
454		Stuck S, Zuber S; SHARE Central Coordination Team (2013) Data resource profile: the
455		survey of health, ageing and retirement in Europe (SHARE). Int J Epidemiol 42, 992-
456		1001.
457	[20]	Freedman VA, Kasper JD (2019) Cohort profile: the National Health and Aging Trends
458		Study (NHATS). Int J Epidemiol 48, 1044-1045g.
459	[21]	Börsch-Supan, A. Survey of Health, Ageing and Retirement in Europe (SHARE) Wave
460		8. Release version: 8.0.0. SHARE-ERIC. Data set,
461		https://doi.org/10.6103/SHARE.w8.800, Last updated February 10, 2022, Accessed on
462		October 30, 2023.
463	[22]	Börsch-Supan A. Survey of Health, Ageing and Retirement in Europe (SHARE) Wave
464		8. COVID-19 Survey 1. Release version: 8.0.0. SHARE-ERIC. Data set,
465		https://doi.org/10.6103/SHARE.w8ca.800, Last updated February 10, 2022, Accessed
466		on October 30, 2023.
467	[23]	Scherpenzeel A, Axt K, Bergmann M, Douhou S, Oepen A, Sand G, Schuller K, Stuck
468		S, Wagner M, Börsch-Supan A (2020) Collecting survey data among the 50+ population
469		during the COVID-19 outbreak: the Survey of Health, Ageing and Retirement in
470		Europe (SHARE). Surv Res Methods 14, 217–221.
471	[24]	Bergmann M. and Börsch-Supan A (Eds.) (2021) SHARE Wave 8 Methodology:

472

Collecting Cross-National Survey Data in Times of COVID-19, Max Planck Institute

- 473 for Social Law and Social Policy, Munich.
- 474 [25] Hofer SM, Piccinin AM (2009) Integrative data analysis through coordination of
- 475 measurement and analysis protocol across independent longitudinal studies. *Psychol*
- 476 *Methods* 14, 150–164.
- 477 [26] Amjad H, Roth DL, Sheehan OC, Lyketsos CG, Wolff JL, Samus QM (2018)
- 478 Underdiagnosis of dementia: an observational study of patterns in diagnosis and
- awareness in US older adults. *J Gen Intern Med* **33**, 1131–1138.
- 480 [27] Lugo-Palacios DG, Gannon B (2017) Health care utilisation amongst older adults with

481 sensory and cognitive impairments in Europe. *Health Econ Rev* 7, 44.

- 482 [28] Sutin AR, Luchetti M, Stephan Y, Terracciano A (2020) Meaning in life and risk of
- 483 cognitive impairment: a 9-year prospective study in 14 countries. *Arch Gerontol*
- 484 *Geriatr* **88**, 104033.
- 485 [29] Luchetti M, Terracciano A, Aschwanden D, Lee JH, Stephan Y, Sutin AR (2020)
- 486 Loneliness is associated with risk of cognitive impairment in the Survey of Health,

487 Ageing and Retirement in Europe. *Int J Geriatr Psychiatry* **35**, 794–801.

488 [30] Mehrbrodt T, Gruber S, Wagner M. SHARE: scales and multi-item indicators,

489 http://www.share-

490 project.org/fileadmin/pdf_documentation/SHARE_Scales_and_Multi-

491		Item_Indicators.pdf. Updated 2019, Last updated January 2021, Accessed October 30,
492		2023.
493	[31]	Kasper JD, Freedman VA, Spillman B. Classification of persons by dementia status in
494		the National Health and Aging Trends Study. NHATS Technical Paper #5,
495		https://www.nhats.org/sites/default/files/2022-
496		09/NHATS%20Dementia%20Classification%20with%20Programming%20Statements_
497		09232022.zip, Last updated April 4, 2023, Accessed October 30, 2023.
498	[32]	Galvin JE, Roe CM, Xiong C, Morris JC (2006) Validity and reliability of the AD8
499		informant interview in dementia. Neurology 67, 1942–1948.
500	[33]	Steffick DE. Documentation of affective functioning measures in the health and
501		retirement study. Ann Arbor: Survey Research Center University of Michigan.
502		https://hrsonline.isr.umich.edu/sitedocs/userg/dr-005.pdf, Last updated 2000, Accessed
503		October 30, 2023.
504	[34]	Prince MJ, Reischies F, Beekman AT, Fuhrer R, Jonker C, Kivela SL, Lawlor BA, Lobo
505		A, Magnusson H, Fichter M, van Oyen H, Roelands M, Skoog I, Turrina C, Copeland
506		JR (1999) Development of the EURO-D scale—a European, union initiative to
507		compare symptoms of depression in 14 European centres. Br J Psychiatry 174, 330-
508		338.

509 [35] Kroenke K, Spitzer RL, Williams JB (2003) The Patient Health Questionnaire-2:

510		validity of a two-item depression screener. Med Care 41, 1284–1292.
511	[36]	Li C, Friedman B, Conwell Y, Fiscella K (2007) Validity of the Patient Health
512		Questionnaire 2 (PHQ-2) in identifying major depression in older people. J Am Geriatr
513		<i>Soc</i> 55 , 596–602.
514	[37]	Di Gessa G, Price D (2021) Changes in health and social well-being in the COVID-19
515		clinically vulnerable older English population during the pandemic. J Epidemiol
516		Community Health 75, 1070–1077.
517	[38]	Steptoe A, Di Gessa G (2021) Mental health and social interactions of older people with
518		physical disabilities in England during the COVID-19 pandemic: a longitudinal cohort
519		study. Lancet Public Health 6, e365–e373.
520	[39]	Zaninotto P, Iob E, Demakakos P, Steptoe A (2022) Immediate and longer-term changes
521		in the mental health and well-being of older adults in England during the COVID-19
522		pandemic. JAMA Psychiatry 79, 151–159.
523	[40]	Wester CT, Bovil T, Scheel-Hincke LL, Ahrenfeldt LJ, Möller S, Andersen-Ranberg K
524		(2022) Longitudinal changes in mental health following the COVID-19 lockdown:
525		Results from the Survey of Health, Ageing, and Retirement in Europe. Ann Epidemiol
526		74, 21–30.
527	[41]	King G, Zeng L (2001) Explaining rare events in international relations. Int Organ 55,
528		693–715.

529	[42]	Van Den Eeckhaut M, Vanwalleghem T, Poesen J, Govers G, Verstraeten G,
530		Vandekerckhove L (2006) Prediction of landslide susceptibility using rare events
531		logistic regression: A case-study in the Flemish Ardennes (Belgium). Geomorphology
532		(Amst) 76 , 392–410.
533	[43]	Cham H, Reshetnyak E, Rosenfeld B, Breitbart W (2017) Full information maximum
534		likelihood estimation for latent variable interactions with incomplete indicators.
535		Multivariate Behav Res 52, 12–30.
536	[44]	Van Winkle Z, Ferragina E, Recchi E (2021) The unexpected decline in feelings of
537		depression among adults ages 50 and older in 11 European countries amid the COVID-
538		19 pandemic. Socius 7, 1–11.
539	[45]	Pierce M, Hope H, Ford T, Hatch S, Hotopf M, John A, Kontopantelis E, Webb R,
540		Wessely S, McManus S, Abel KM (2020) Mental health before and during the COVID-
541		19 pandemic: a longitudinal probability sample survey of the UK population. Lancet
542		Psychiatry 7, 883–892.
543	[46]	Santini ZI, Jose PE, York Cornwell E, Koyanagi A, Nielsen L, Hinrichsen C, Meilstrup
544		C, Madsen KR, Koushede V (2020) Social disconnectedness, perceived isolation, and
545		symptoms of depression and anxiety among older Americans (NSHAP): a longitudinal
546		mediation analysis. Lancet Public Health 5, e62–70.
547	[47]	Talbot CV, Briggs P (2021) 'Getting back to normality seems as big of a step as going

548	into lockdown': the impact of the COVID-19 pandemic on people with early to middle
549	stage dementia. Age Ageing 50, 657–663.

- 550 [48] Burton A, Rapaport P, Palomo M, Lord K, Budgett J, Barber J, Hunter R, Butler L,
- 551 Vickerstaff J, Rockwood K, Ogden M, Smith D, Lang I, Livingston G, Dow B, Kales
- 552 H, Manthorpe J, Walters K, Hoe J, Orgeta V, Samus Q, Cooper C; NIDUS study team
- 553 (2021) Clinical and cost-effectiveness of a New psychosocial intervention to support
- 554 Independence in Dementia (NIDUS-family) for family carers and people living with
- dementia in their own homes: a randomised controlled trial. *Trials* **22**, 865.
- 556 [49] Cooper C, Mansour H, Carter C, Rapaport P, Morgan-Trimmer S, Marchant NL, Poppe
- 557 M, Higgs P, Brierley J, Solomon N, Budgett J, Bird M, Walters K, Barber J, Wenborn J,
- 558 Lang IA, Huntley J, Ritchie K, Kales HC, Brodaty H, Aguirre E, Betz A, Palomo M
- 559 (2021) Social connectedness and dementia prevention: pilot of the APPLE-Tree video-
- call intervention during the Covid-19 pandemic. *Dementia (London)* **20**, 2779–2801.
- 561 [50] Kelleher D, Lord K, Duffy L, Rapaport P, Barber J, Manthorpe J, Leverton M, Dow B,
- 562 Budgett J, Banks S, Duggan S, Cooper C (2022) Time to reflect is a rare and valued
- 563 opportunity; a pilot of the NIDUS-professional dementia training intervention for
- 564 homecare workers during the Covid-19 pandemic. *Health Soc Care Community* **30**,
- 565 e2928–e2939.



567		Winbolt M, Clarke P, Burton J, Low LF, Loi SM, Fairhall A, Polacsek M, Stiles J,
568		Muliadi F, Chau N, Scherer S, Ames D, Sousa TV, Dow B (2021) Promoting
569		Independence Through quality dementia Care at Home (PITCH): a research protocol
570		for a stepped-wedge cluster-randomised controlled trial. Trials 22, 949.
571	[52]	Freedman VA, Hu M, Kasper JD (2022) Changes in older adults' social contact during
572		the COVID-19 pandemic. J Gerontol B Psychol Sci Soc Sci 77, e160–e166.
573	[53]	Comas-Herrera A, Glanz A, Curry N, Deeny S, Hatton C, Hemmings N, Humphries R,
574		Lorenz-Dant K, Oung C, Rajan S, Suarez-Gonzalez A. The COVID-19 Long-Term
575		Care situation in England, https://ltccovid.org/wp-content/uploads/2020/11/COVID-19-
576		Long-Term-Care-situation-in-England-19-November-2.pdf, Last updated November 19,
577		2020, Accessed October 30, 2023.
578	[54]	Sriram V, Jenkinson C, Peters M (2021) Impact of COVID-19 restrictions on carers of
579		persons with dementia in the UK: a qualitative study. Age Ageing 50, 1876–1885.
580	[55]	Vislapuu M, Angeles RC, Berge LI, Kjerstad E, Gedde MH, Husebo BS (2021) The
581		consequences of COVID-19 lockdown for formal and informal resource utilization
582		among home-dwelling people with dementia: results from the prospective PAN.DEM
583		study. BMC Health Serv Res 21, 1003.
584	[56]	Dyer AH, Murphy C, Lawlor B, Kennelly SP, For The Nilvad Study Group (2021)
585		Social networks in mild-to-moderate Alzheimer disease: longitudinal relationships with

587		1923–1929.
588	[57]	Bakker ED, van Maurik IS, Mank A, Zwan MD, Waterink L, van den Buuse S, van den
589		Broeke JR, Gillissen F, van de Beek M, Lemstra E, van den Bosch KA, van
590		Leeuwenstijn M, Bouwman FH, Scheltens P, van der Flier WM (2022) Psychosocial
591		effects of COVID-19 measures on (pre-)dementia patients during second lockdown. J
592		Alzheimers Dis 86, 931–939.
593	[58]	Clare L, Martyr A, Gamble LD, Pentecost C, Collins R, Dawson E, Hunt A, Parker S,
594		Allan L, Burns A, Hillman A, Litherland R, Quinn C, Matthews FE, Victor C (2022)
595		Impact of COVID-19 on 'living well' with mild-to-moderate dementia in the
596		community: findings from the IDEAL cohort. J Alzheimers Dis 85, 925-940.

dementia severity, cognitive function, and adverse events. Aging Ment Health 25,

597 List of Supplementary Materials

- **598 Table S1.** Variables across three cohort studies
- **Table S2.** Presence of depressive symptoms according to presence of dementia at baseline
- across England, other 27 European countries, and the United States
- **Table S3.** Coefficients, 95% confidence intervals, and P-values of depressive symptoms of all
- 602 covariates in the first model across England, other 27 European countries, and the United
- 603 States
- **Table S4.** Sensitivity analysis of the first model replacing with four categories of home-care
- use: coefficients, 95% confidence intervals, and P-values of depressive symptoms of all
- 606 covariates across England and the United States
- **Table S5.** Sensitivity analysis of the first model excluding participants reduced home-care
- use: coefficients, 95% confidence intervals, and P-values of depressive symptoms of all
- 609 covariates across England, other 27 European countries, and the United States

Table S6. Coefficients, 95% confidence intervals, and P-values of depressive symptoms of all

611 covariates in the second model across England, other 27 European countries, and the United

612 States

- **Table S7.** Sensitivity analysis of the second model replacing with total scores of depressive
- 614 symptoms: coefficients, 95% confidence intervals, and P-values of depressive symptoms of
- all covariates across England and the United States

	ELSA		SHARE		NHATS	
	N of responses	Mean (SD) or N	N of responses	Mean (SD) or N	N of responses	Mean (SD) or N
		(%)		(%)		(%)
Age, mean (SD)			34,475	70.4 (9.1)	3,001	79.6 (6.5)
Age, N (%)	6,306		34,475		3,001	
74 or less		2,587 (41.0)		13,394 (38.9)		738 (24.6)
75–84		1,162 (18.4)		8,530 (24.7)		1,561 (52.0)
85 or more		182 (2.9)		2,557 (7.4)		702 (23.4)
Sex, man, N (%)	6,306	2,772 (44.0)	34,475	14,510 (42.1)	3,001	1,290 (43.0)
Married or with a	6,305	4,348 (69.0)	34,475	23,735 (68.8)	3,001	1,542 (51.4)

Table 1. Characteristics of older adults in the three cohort studies prior to the COVID-19 pandemic

partner, N (%)

Living alone, N (%)	6,306	1,264(20.0)	34,475	10,515 (30.5)	3,001	934 (31.1)
Educational attainment,	5,747		34,475		2,965	
N (%)						
1 (low)		840 (14.6)		5,910 (17.1)		473 (15.8)
2		1,603 (27.9)		5,741 (16.7)		764 (25.8)
3		733 (12.8)		13,104 (38.0)		823 (27.8)
4 (high)		2,571 (44.7)		9,720 (28.2)		941 (31.4)
Dementia, N (%)	6,114	179 (2.9)	33,623	773 (2.3)	3,001	194 (6.5)

Note: COVID-19, coronavirus disease 2019; ELSA, English Longitudinal Study of Ageing (N = 6,306); SHARE, Study of Health, Ageing and

618 Retirement in Europe (N = 34,475); NHATS, National Health and Aging Trends Study (N = 3,001); SD, standard deviation; N, number.

	ELSA		SHARE		NHATS		
	Dementia	No dementia	Dementia	No dementia	Dementia	No dementia	
Home-care use, N (%)	N = 178	N = 5,934	N = 772	N = 32,824	N = 182	N = 2,677	
Reduced	3 (1.7)	30 (0.5)	22 (2.8)	408 (1.2)	2 (1.1)	24 (0.9)	
Same or increased	27 (15.2)	360 (6.1)	98 (12.7)	1,396 (4.3)	142 (78.0)	1,230 (45.9)	
Same	16 (9.0)	263 (4.4)	N/A	N/A	128 (70.3)	1,109 (41.4)	
Increased	11 (6.2)	97 (1.6)	N/A	N/A	14 (7.7)	121 (4.5)	
No home care used	148 (83.1)	5,544 (93.4)	652 (84.5)	31,020 (94.5)	38 (20.9)	1,423 (53.1)	

619 ′	Table 2. Change in home-care	use during the COVID-19	pandemic by dementia at baseline
-------	------------------------------	-------------------------	----------------------------------

620 *Note:* COVID-19, coronavirus disease 2019; ELSA, English Longitudinal Study of Ageing (N = 6,112; 194 excluded due to missing data);

621 SHARE, Study of Health, Ageing and Retirement in Europe (N = 33,596; 879 excluded due to missing data); NHATS, National Health and

622 Aging Trends Study (N = 2,859; 142 excluded due to missing data). In SHARE, positive response (yes) to the question whether facing more

- 623 difficulties in getting the amount of home care needed was coded as "Reduced." Negative response (no) to the question was coded as "Same or
- 624 increased."

Table 3. Odds ratios, 95% confidence intervals, and P-values for the presence of depressive symptoms by baseline dementia and change in

	ELSA		SHARE		NHATS	
	Odds ratio (95%CI)	P-value	Odds ratio (95%CI)	P-value	Odds ratio (95%CI)	P-value
Within-person level						
Home care during the pandemic,						
reference = no home care used						
Reduced	11.236 (3.230, 39.084)	<.001	2.444 (1.897, 3.149)	<.001	5.401 (1.282, 22.751)	.022
Same or increased	2.974 (2.127, 4.158)	<.001	1.681 (1.456, 1.940)	<.001	2.305 (1.681, 3.162)	<.001
Period, during the pandemic	3.695 (3.264, 4.185)	<.001	0.375 (0.360, 0.391)	<.001	1.157 (0.885, 1.512)	.285
Dementia x during pandemic	0.499 (0.304, 0.819)	.006	0.809 (0.635, 1.030)	.086	0.890 (0.483, 1.640)	.709
Between-person level	Coefficient (95%CI)		Coefficient (95%CI)		Coefficient (95%CI)	

626 home-care use during the COVID-19 pandemic

	Dementia at baseline	1.531 (1.107, 1.954)	<.001	0.882 (0.701, 1.063)	<.001	1.474 (1.007, 1.865)	<.001
627	Note: Multilevel binomial logisti	c regression analysis was em	ployed. All	analyses were adjusted for	age, sex, m	narried or with a partner, li	iving
628	alone, and educational attainmen	t. Full-information maximun	n likelihood	methods were used to hand	dle missing	data. CI, confidence inter	val.
629	ELSA, English Longitudinal Stud	dy of Ageing (N = $6,306$); De	epressive sy	mptoms were measured us	ing a shorte	ned version of the Center	for
630	Epidemiologic Studies Depressio	n Scale (total score ≥4). SHA	ARE, Study	of Health, Ageing and Ret	irement in I	Europe (N = 34,475); Pres	ence of
631	depressive symptoms was assessed	ed by single question for dep	ression from	n European-Depression sca	ile: "In the l	ast month, have you been	sad or
632	depressed?" NHATS, National H	ealth and Aging Trends Stud	y (N = 3,00	1); Presence of depressive	symptoms v	vere measured using the P	atient
633	Health Questionnaire-2 (total sco	re ≥3).					

	ELSA		SHARE		NHATS	
	Odds ratio (95%CI)	P-value	Odds ratio (95%CI)	P-value	Odds ratio (95%CI)	P-value
Within-person level						
Home care during the						
pandemic, reference = no						
home care used						
Reduced	12.798 (3.627, 45.155)	<.001	2.505 (1.938, 3.237)	<.001	4.111 (1.047, 16.136)	.043
Same or increased	3.100 (2.188, 4.393)	<.001	1.712 (1.477, 1.984)	<.001	2.322 (1.704, 3.164)	<.001
Period, during the	3.601 (3.185, 4.071)	<.001	0.378 (0.362, 0.394)	<.001	1.116 (0.859, 1.449)	.410

Table 4. Odds ratios, 95% confidence intervals, and P-values for the presence of depressive symptoms by interaction between baseline dementia

and change in home-care use during the COVID-19 pandemic

pandemic

Dementia x during	0.690 (0.413, 1.151)	.155	0.842 (0.651, 1.090)	.192	1.114 (0.458, 2.708)	.812
pandemic						
Dementia x reduced	0.202 (0.004, 9.817)	.419	0.720 (0.114, 4.532)	.726	999.000 (999.000,	.187
					999.000)	
Dementia x same or	0.690 (0.209, 2.278)	.543	0.722 (0.395, 1.322)	.291	0.718 (0.261, 1.973)	.521
increased						
Between-person level	Coefficient (95%CI)		Coefficient (95%CI)		Coefficient (95%CI)	
Dementia at baseline	1.265 (0.872, 1.658)	<.001	0.893 (0.701, 1.084)	<.001	1.389 (0.976, 1.803)	<.001

636 Note: Multilevel binomial logistic regression analysis was employed. All analyses were adjusted for age, sex, married or with a partner, living

637 alone, and educational attainment. Full-information maximum likelihood methods were used to handle missing data. CI, confidence interval.

ELSA, English Longitudinal Study of Ageing (N = 6,306); Depressive symptoms were measured using a shortened version of the Center for

639 Epidemiologic Studies Depression Scale (total score \geq 4). SHARE, Study of Health, Ageing and Retirement in Europe (N = 34,475); Presence of

641 depressed?" NHATS, National Health and Aging Trends Study (N = 3,001); Presence of depressive symptoms were measured using the Patient

642 Health Questionnaire-2 (total score \geq 3).

Figure Legends

Figure 1. Flow chart of this study

Numbers of participants in COVID-19 surveys in this figure may not be equal to those in the original data sets, as some participants were excluded based on the exclusion criteria for this study. ELSA, English Longitudinal Study of Ageing; SHARE, Study of Health, Ageing and Retirement in Europe; EU, European Union; NHATS, National Health and Aging Trends Study.

Figure 2. Presence of depressive symptoms before and during the COVID-19 pandemic across England, other European countries, and the United States

Baseline, main wave survey before the COVID-19 pandemic. Follow-up, sub-study during the pandemic. A: ELSA, English Longitudinal Study of Ageing. Depressive symptoms were measured using a shortened version of the Center for Epidemiologic Studies Depression Scale (total score \geq 4). B: SHARE, Study of Health, Ageing and Retirement in Europe. Presence of depressive symptoms was assessed by single question for depression from European-Depression scale: "In the last month, have you been sad or depressed?" C: NHATS, National Health and Aging Trends Study. Presence of depressive symptoms were measured using the Patient Health Questionnaire-2 (total score \geq 3).

Table S1. Variables across the three cohort studies

	Conort		
	ELSA	SHARE	NHATS
Variable	England	27 European countries	The United States
 Depressive symptoms Dementia status Age 	CESD score, range 0–8	One item from EURO-D: "In the last	PHQ-2 score, range 0–6
symptoms	• 1 = present (4 or more)	month, have you been sad or	 1 = present (3 or more)
	• 0 = no (0–3)	depressed?"	• 0 = no (0–2)
		• 1 = yes	
		• 0 = no	
Dementia status	Classification approach using	Classification approach using	Dementia classification by validated
	immediate recall, delayed recall, and	immediate recall, delayed recall, and	algorithm for survey
	verbal fluency	verbal fluency	
	• 1 = dementia	• 1 = dementia	• 1 = dementia
	• 0 = no dementia	• 0 = no dementia	• 0 = no dementia
Age	Original cohort sampled from adults	Original cohort sampled from adults	Original cohort sampled from adults
	aged 50-89 or "90 or more" (classified	aged 50 or more	aged 65 or more
	into one category)		
	Reclassified into five age bands	Reclassified into five age bands	Reclassified into five age bands
	• 1 = 74 or younger	• 1 = 74 or younger	• 1 = 74 or younger
	• 2 = 75-84	• 2 = 75-84	• 2 = 75-84
	• 3 = 85 or older	• 3 = 85 or older	• 3 = 85 or older
Sex	Male or female	Male or female	Male or female
Educational	• 1 = NVQ4/NVQ5/degree or	ISCED-97	• 1 = no schooling completed
	Variable Depressive symptoms Dementia status Age Sex Educational	ELSAVariableEnglandDepressiveCESD score, range 0–8symptoms• 1 = present (4 or more)• 0 = no (0–3)Dementia statusClassification approach using immediate recall, delayed recall, and verbal fluency• 1 = dementia• 0 = no dementiaAgeOriginal cohort sampled from adults aged 50-89 or "90 or more" (classified 	ELSASHAREVariableEngland27 European countriesDepressiveCESD score, range 0–8One item from EURO-D: "In the lastsymptoms• 1 = present (4 or more)month, have you been sad or• 0 = no (0-3)depressed?"• 1 = yes• 0 = noDementia statusClassification approach using immediate recall, delayed recall, and verbal fluencyClassification approach using immediate recall, delayed recall, and verbal fluency• 1 = dementia• 1 = dementia• 0 = no dementia• 0 = no dementiaAgeOriginal cohort sampled from adults aged 50-89 or "90 or more" (classified into one category)Original cohort sampled from adults aged 50 or moreReclassified into five age bands • 1 = 74 or younger • 2 = 75-84 • 3 = 85 or olderReclassified into five age bands • 1 = 74 or younger • 2 = 75-84 • 3 = 85 or olderSexMale or femaleMale or femaleEducational• 1 = NVQ4/NVQ5/degree orISCED-97

		Cohort		
		ELSA	SHARE	NHATS
Time	Variable	England	27 European countries	The United States
	attainment	equivalent	• 0 = pre-primary education	• 2 = 1st–8th grade
		• 2 = higher education below	• 1 = primary or first stage of basic	• 3 = 9th–12th grade
		degree	education	• 4 = high school graduate (high
		• 3 = NVQ3/GCE advanced level	• 2 = lower secondary or second	school diploma or equivalent)
		equivalent	stage of basic education	• 5 = vocational, technical,
		• 4 = NVQ2/GCE ordinary level	• 3 = (upper) secondary education	business, or trade school
		equivalent	• 4 = post-secondary non-tertiary	certificate or diploma (beyond
		• 5 = NVQ1/CSE other grade	education	high school)
		equivalent	• 5 = first stage of tertiary	• 6 = some college but no degree
		• 6 = foreign/other	education	• 7 = associate's degree
		• 7 = no qualification	• 6 = second stage of tertiary	• 8 = bachelor's degree
			education	• 9 = master's, professional, or
				doctoral degree
		Reclassification using quartiles	Reclassification using quartiles	Reclassification using quartiles
		• 1 = no qualification	• 1 = ISCED 97 level 0 OR 1	• 1 = no school completed OR 1st-
		• 2 = NVQ2/GCE ordinary level OR	• 2 = ISCED 97 level 2	8th grade OR 9th–12th grade
		NVQ1/CSE	• 3 = ISCED level 3	• 2 = high school graduate
		• 3 = NVQ3/GCE advanced level	• 4 = ISCED level 4 OR 5 OR 6	• 3 = diploma beyond high school
		OR higher education below		OR some college OR associate's
		degree		degree

		Cohort		
		ELSA	SHARE	NHATS
Time	Variable	England	27 European countries	The United States
		4 = NVQ4/NVQ5missing = foreign/other		 4 = bachelor's OR master's, professional, or doctoral degree
	Married or with a partner	 1 = yes: married OR remarried 0 = no: single, never married OR separated OR divorced OR widowed 	 1 = yes: married and living together OR registered partnership OR married, living separated 0 = never married OR divorced OR widowed 	 1 = yes: married OR living with a partner 0 = no: separated OR divorced OR widowed OR never married
	Living alone	 1 = yes: number of people in household is equal to 1 0 = no: number of people in household >1 	 1 = yes: household type is single person 0 = no: household type is other than single person 	 1 = yes: number of people in household is equal to 1 0 = no: no: number of people in household >1
Follow-up	Depressive symptoms	 CESD score, range 0–8 1 = present (4 or more) 0 = no (0–3) 	One item from EURO-D: "In the last month, have you been sad or depressed?" • 1 = yes • 0 = no	 PHQ-2 score, range 0–6 1 = present (3 or more) 0 = no (0–2)
	Home-care use	Over the past month have you received care at home? • Yes, formal (paid, provided from an agency)	Did you regularly receive home carebefore the outbreak of Corona?YesNo	 DURING the COVID-19 outbreak, in a typical week, how many people have done household activities with you or

		Cohort				
		ELSA	SHARE	NHATS		
Time	Variable	England	27 European countries	The United States		
		• Yes, informal (friend or relative)		for you or helped you with		
		• No		personal care activities?		
				DURING the COVID-19		
				outbreak, in a typical week,		
				about how many hours have		
				people spent doing your		
				household activities with you o		
				for you or helping you with		
				personal care activities?		
	Change in home-	Since the coronavirus outbreak	Since the outbreak of Corona, did you	Is that more, less or about the same		
	care use	started is the amount of care you	face more difficulties in getting the	compared to a typical week before		
		receive	amount of home care that you need?	the COVID-19 outbreak?		
		Less than it was	• Yes	Less than before		
		About the same	• No	More than before		
		• More than it was		About the same		
		No longer receive help				
	Reclassification of	"No home care use": Did not	"No home care use": Did not	• "No home care use": No peopl		
	change in home-	receive care at home over the	regularly receive home care	helped in a typical week		
	care use	past month OR no longer receive	• "Same or increased": Did not	"Same or increased": Hours		
		help	face more difficulties in getting	people spent were about the		
		• "Same or increased": the amount	the amount of home care	same OR more than before		

		Cohort			
	ELSA		SHARE	NHATS	
Time Variable		England	27 European countries	The United States	
		of care was same OR more than	needed	"Reduced": Hours people spent	
		it was	"Reduced": Faced more	were less than before	
		• "Reduced": the amount of care	difficulties		
		was less than it was			

Note: ELSA, English Longitudinal Study of Ageing (N = 6,306); SHARE, Study of Health, Ageing and Retirement in Europe (N = 34,475); NHATS, National Health and Aging Trends Study (N =

3,001); CESD, a shortened version of the Center for Epidemiologic Studies Depression Scale; EURO-D, European-Depression scale; PHQ-2, Patient Health Questionnaire-2; NVQ, National

Vocational Qualification; GCE, General Certificate of Education; CSE, Certificate of Secondary Education; ISCED, International Standard Classification of Education.

•								
N (%)	ELSA		SHARE		NHATS	NHATS		
	Dementia	No dementia	Dementia	No dementia	Dementia	No dementia		
Baseline								
Depressive symptoms								
Yes	54 (31.8)	641 (10.9)	425 (57.1)	12,770 (38.9)	42 (22.5)	203 (7.3)		
No	116 (68.2)	5,259 (89.1)	319 (42.9)	20,040 (61.1)	145 (77.5)	2596 (92.7)		
Follow-up								
Depressive symptoms								
Yes	71 (40.1)	1,429 (24.4)	287 (37.5)	8,158 (24.9)	59 (31.4)	300 (10.8)		
No	106 (59.9)	4,428 (75.6)	478 (62.5)	24,620 (75.1)	129 (68.6)	2488 (89.2)		

Table S2. Presence of depressive symptoms according to presence of dementia at baseline across England, other 27 European countries, and the United States

Variable	ELSA			SHARE			NHATS		
	Coefficient	95%CI	P-value	Coefficient	95%CI	P-value	Coefficient	95%CI	P-value
Within-person level									
Home care during the									
pandemic, reference =									
no home care used									
Reduced	2.419	1.172, 3.666	<.001	0.894	0.640, 1.147	<.001	1.687	0.249, 3.125	.022
Same or increased	1.090	0.755, 1.425	<.001	0.519	0.376, 0.663	<.001	0.835	0.519, 1.151	<.001
Period, during the	1 207	1 102 1 121	< 001	-0.980	-1.022, -0.938	<.001	0.146	-0.122, 0.414	.285
pandemic	1.307	1.100, 1.401 5.001							
Dementia x during	0.605	1 1 2 0 0 2 0 0	006	-0.212	-0.454, 0.030	.086	-0.116	-0.727, 0.495	.709
pandemic	-0.095	-1.169, -0.200	.000						
Between-person level									
Dementia at baseline	1.531	1.107, 1.954	<.001	0.882	0.701, 1.063	<.001	1.474	1.007, 1.940	<.001
Age, reference = 74 or									
less									
75-84	-0.402	-0.608, -0.197	<.001	0.259	0.197, 0.320	<.001	0.039	-0.349, 0.427	.843
85 or more	-0.539	-1.034, -0.044	.033	0.314	0.212, 0.416	<.001	-0.058	-0.496, 0.380	.795
Sex, male	-0.770	-0.928, -0.612	<.001	-0.853	-0.907, -0.799	<.001	-0.028	-0.318, 0.261	.849
Married or with a partner	-0.517	-0.762, -0.272	<.001	-0.011	-0.123, 0.102	.851	-0.466	-0.797, -0.135	.006
Living alone	0.463	0.195, 0.731	<.001	0.464	0.352, 0.576	<.001	0.053	-0.282, 0.387	.757
Educational attainment,									

Table S3. Coefficients, 95% confidence intervals, and P-values of depressive symptoms of all covariates in the first model across England, other 27 European countries, and the United States

reference = 1 (low)

2	-0.387	-0.646, -0.129	.003	-0.143	-0.233, -0.053	.002	-0.432	-0.823, -0.041	.030
3	-0.371	-0.697, -0.045	.026	-0.257	-0.335, -0.179	<.001	-0.999	-1.424, -0.573	<.001
4 (high)	-0.509	-0.755, -0.264	<.001	-0.362	-0.444, -0.280	<.001	-1.521	-1.960, -1.082	<.001

Note: Multilevel binomial logistic regression analysis was employed. Full-information maximum likelihood methods were used to handle missing data. CI, confidence interval. ELSA, English

Longitudinal Study of Ageing (N = 6,306); SHARE, Study of Health, Ageing and Retirement in Europe (N = 34,475); NHATS, National Health and Aging Trends Study (N = 3,001).

Variable	ELSA			NHATS		
	Coefficient	95%CI	P-value	Coefficient	95%CI	P-value
Within-person level						
Home care during the pandemic,						
reference = no home care used						
Reduced	2991.107	2950.957, 3031.257	<.001	1.701	0.398, 3.004	.011
Same	3567.569	3528.318, 3606.819	<.001	0.717	0.398, 1.037	<.001
Increased	-538.428	-538.428, -538.428	.999	1.529	0.908, 2.150	<.001
Period, during the pandemic	-13710.145	-13867.497, -13552.792	<.001	0.141	-0.120, 0.403	.290
Dementia x during pandemic	62.470	-109.125, 234.065	.476	-0.048	-0.652, 0.556	.876
Between-person level						
Dementia at baseline	1.479	1.069, 1.888	<.001	1.374	0.914, 1.834	<.001
Age, reference = 74 or less						
75-84	-0.354	-0.640, -0.068	<.001	-0.114	-0.436, 0.208	.489
85 or more	-0.267	-0.927, 0.393	.015	-0.221	-0.625, 0.183	.283
Sex, male	-0.686	-0.925, -0.447	.428	0.004	-0.282, 0.291	.976
Married or with a partner	-0.844	-1.168, -0.521	<.001	-0.519	-0.863, -0.176	.003
Living alone	0.392	0.064, 0.720	.019	-0.022	-0.334, 0.291	.891
Educational attainment, reference						
= 1 (low)						
2	-0.670	-1.030, -0.309	<.001	-0.504	-0.880, -0.129	.009
3	-0.970	-1.414, -0.526	<.001	-1.108	-1.521, -0.695	<.001

Table S4. Sensitivity analysis of the first model replacing with four categories of home-care use: coefficients, 95% confidence intervals, and P-values of depressive symptoms of all covariates across England and the United States

Note: Multilevel binomial logistic regression analysis was employed. Full-information maximum likelihood methods were used to handle missing data. CI, confidence interval. ELSA, English Longitudinal Study of Ageing (N = 6,306); NHATS, National Health and Aging Trends Study (N = 3,001).

Table S5. Sensitivity analysis of the first model excluding participants reduced home-care use: coefficients, 95% confidence intervals, and P-values of depressive symptoms of all covariates across England, other 27 European countries, and the United States

Variable	ELSA			SHARE			NHATS		
	Coefficient	95%CI	P-value	Coefficient	95%CI	P-value	Coefficient	95%CI	P-value
Within-person level									
Home care during the pandemic,									
reference = no home care used									
Same or increased				0.524	0.380, 0.667	<.001			
Same	E40.040	533.869,	< 001				0.746	0.419, 1.073	<.001
	549.818	565.767	<.001						
Increased	204 222	-294.322, -	000				1.543	0.902, 2.184	<.001
	-294.322	294.322	.999						
Period, during the pandemic	1072 000	-1873.888, -	000	-0.967	-1.009, -0.925	<.001	0.156	-0.114, 0.426	.257
	-10/3.000	1873.888	.999						
Dementia x during pandemic	116 005	-2403.642,	020	-0.228	-0.481, 0.026	.078	-0.072	-0.685, 0.540	.817
	110.000	2687.411	.928						
Between-person level									
Dementia at baseline	0.854	0.607, 1.100	<.001	0.898	0.699, 1.096	<.001	1.389	0.915, 1.863	<.001
Age, reference = 74 or less									
75-84	-0.291	-0.525, -0.057	.015	0.257	0.195, 0.319	<.001	0.007	-0.347, 0.360	.971
85 or more	-0.116	-0.529, 0.298	.584	0.303	0.199, 0.408	<.001	-0.055	-0.485, 0.374	.801
Sex, male	-0.587	-0.799, -0.376	<.001	-0.857	-0.911, -0.803	<.001	-0.017	-0.301, 0.268	.909
Married or with a partner	-0.801	-1.036, -0.565	<.001	-0.001	-0.107, 0.106	.990	-0.451	-0.804, -0.098	.012

Living a	alone	0.147	-0.034, 0.328	.112	0.472	0.364, 0.579	<.001	0.068	-0.268, 0.405	.690
Educat	ional attainment, reference									
= 1 (lov	v)									
2		-0.446	-0.655, -0.237	<.001	-0.149	-0.237, -0.061	.001	-0.481	-0.850, -0.113	.010
3		-0.703	-1.038, -0.368	<.001	-0.253	-0.331, -0.176	<.001	-1.094	-1.482, -0.706	<.001
4	(high)	-0.606	-0.830, -0.381	<.001	-0.360	-0.440, -0.279	<.001	-1.608	-2.008, -1.209	<.001

60

Note: Multilevel binomial logistic regression analysis was employed. Full-information maximum likelihood methods were used to handle missing data. CI, confidence interval. ELSA, English

Longitudinal Study of Ageing (N = 6,273); SHARE, Study of Health, Ageing and Retirement in Europe (N = 34,018); NHATS, National Health and Aging Trends Study (N = 2,975).

Variable	ELSA			SHARE			NHATS		
	Coefficient	95%CI	P-value	Coefficient	95%CI	P-value	Coefficient	95%CI	P-value
Within-person									
level									
Home care during									
the pandemic,									
reference = no									
home care used									
Reduced	2.549	1.288, 3.810	<.001	0.918	0.661, 1.175	<.001	1.414	0.046, 2.781	.043
Same or	1.132	0.783, 1.480	<.001	0.538	0.390, 0.685	<.001	0.842	0.533, 1.152	<.001
increased									
Period, during the	1.281	1.158, 1.404	<.001	-0.974	-1.015, -0.933	<.001	0.110	-0.151, 0.371	.410
pandemic									
Dementia x during	0.271	-1.565, 0.823	.155	-0.172	-0.430, 0.086	.192	0.108	-0.781, 0.996	.812
pandemic	-0.371								
Dementia x	-1.601	-5.487, 2.284	.419	-0.329	-2.169, 1.511	.726	47.305	-22.933, 117.548	.187
reduced									
Dementia x same	-0.371	-1.565, 0.823	.543	-0.325	-0.930, 0.279	.291	-0.331	-1.343, 0.680	.521
or increased									
Between-person									
level									
Dementia at	1.265	0.872 1.658	<.001	0.893	0.701, 1.084	<.001	1.389	0.976, 1.803	<.001

Table S6. Coefficients, 95% confidence intervals, and P-values of depressive symptoms of all covariates in the second model across England, other 27 European countries, and the United States

baseline									
Age, reference =									
74 or less									
75-84	-0.412	-0.622, -0.201	.001	0.261	0.200, 0.321	<.001	0.045	-0.264, 0.353	.777
85 or more	-0.589	-1.014, -0.164	.007	0.325	0.224, 0.423	<.001	-0.030	-0.363, 0.304	.861
Sex, male	-0.778	-0.938, -0.618	<.001	-0.839	-0.893, -0.786	<.001	0.040	-0.235, 0.314	.777
Married or with a partner	-0.670	-0.882, -0.459	<.001	0.026	-0.076, 0.128	.616	-0.364	-0.692, -0.036	.029
Living alone	0.275	0.040, 0.510	.022	0.487	0.385, 0.590	<.001	0.142	-0.211, 0.496	.430
Educational									
attainment,									
reference = 1									
(low)									
2	-0.466	-0.704, -0.227	<.001	-0.138	-0.223, -0.052	.002	-0.370	-0.726, -0.013	.042
3	-0.419	-0.726, -0.112	.007	-0.240	-0.316, -0.164	<.001	-0.917	-1.308, -0.527	<.001
4 (high)	-0.585	-0.813, -0.357	<.001	-0.337	-0.416, -0.258	<.001	-1.428	-1.794, -1.063	<.001

Note: Multilevel binomial logistic regression analysis was employed. Full-information maximum likelihood methods were used to handle missing data. CI, confidence interval. ELSA, English Longitudinal Study of Ageing (N = 6,306); SHARE, Study of Health, Ageing and Retirement in Europe (N = 34,475); NHATS, National Health and Aging Trends Study (N = 3,001).

2 of depressive symptoms: coefficients, 95% confidence intervals, and P-values

3 of depressive symptoms of all covariates across England and the United

4 States

Variable	ELSA			NHATS		
	Coefficient	95%CI	P-value	Coefficient	95%CI	P-'
Within-person level						
Home care during the						
pandemic, reference = no home						
care used						
Reduced	1.891	1.085, 2.698	<.001	0.664	0.051, 1.277	.03
Same or increased	1.172	0.919, 1.426	<.001	0.233	0.149, 0.317	<.(
Period, during the pandemic	0.106	0.051, 0.162	<.001	0.074	0.019, 0.130	.00
Dementia x during pandemic	0.062	-0.344, 0.469	.765	0.164	-0.233, 0.561	.41
Dementia x reduced	-1.091	-3.753, 1.572	.422	0.131	-1.927, 2.189	.90
Dementia x same or increased	0.082	-0.936, 1.099	.875	-0.153	-0.626, 0.321	.52
Between-person level						
Dementia at baseline	0.691	0.413, 0.969	<.001	0.627	0.389, 0.866	<.(
Age, reference = 74 or less						
75-84	-0.221	-0.319, -0.123	<.001	0.003	-0.087, 0.094	.94
85 or more	-0.347	-0.541, -0.154	<.001	-0.012	-0.125, 0.101	.83
Sex, male	-0.474	-0.551, -0.397	<.001	-0.130	-0.209, -0.050	.00
Married or with a partner	-0.262	-0.390, -0.135	<.001	-0.097	-0.199, 0.005	.06
Living alone	0.309	0.151, 0.466	<.001	0.065	-0.039, 0.170	.22
Educational attainment,						
reference = 1 (low)						
2	-0.172	-0.312, -0.032	.016	-0.242	-0.382, -0.101	.00
3	-0.228	-0.389, -0.067	.006	-0.446	-0.582, -0.310	<.(
4 (high)	-0.270	-0.400, -0.140	<.001	-0.567	-0.699, -0.435	<.(

5 Note: Multilevel linear regression analysis was employed. Full-information maximum likelihood methods were used to handle

6 missing data. CI, confidence interval. ELSA, English Longitudinal Study of Ageing (N = 6,306); NHATS, National Health and

7 Aging Trends Study (N = 3,001).

8

9