



# The Impact of COVID-19 Quarantine on Patients With Dementia and Family Caregivers: A Nation-Wide Survey

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**Introduction:** Previous studies showed that quarantine for pandemic diseases is associated with several psychological and medical effects. The consequences of quarantine for COVID-19 pandemic in patients with dementia are unknown. We investigated the clinical changes in patients with Alzheimer's disease and other dementias, and evaluated caregivers' distress during COVID-19 quarantine.

**Methods:** The study involved 87 Italian Dementia Centers. Patients with Alzheimer's Disease (AD), Dementia with Lewy Bodies (DLB), Frontotemporal Dementia (FTD), and Vascular Dementia (VD) were eligible for the study. Family caregivers of patients with dementia were interviewed by phone in April 2020, 45 days after quarantine declaration. Main outcomes were patients' changes in cognitive, behavioral, and motor symptoms. Secondary outcomes were effects on caregivers' psychological features.

**Results:** 4913 patients (2934 females, 1979 males) fulfilled the inclusion criteria. Caregivers reported a worsening in cognitive functions in 55.1% of patients, mainly in subjects with DLB and AD. Aggravation of behavioral symptoms was observed in 51.9% of patients. In logistic regression analysis, previous physical independence was

associated with both cognitive and behavioral worsening (odds ratio 1.85 [95% CI 1.42–2.39], 1.84 [95% CI 1.43–2.38], respectively). On the contrary, pandemic awareness was a protective factor for the worsening of cognitive and behavioral symptoms (odds ratio 0.74 [95% CI 0.65–0.85]; and 0.72 [95% CI 0.63–0.82], respectively). Approximately 25.9% of patients showed the onset of new behavioral symptoms. A worsening in motor function was reported by 36.7% of patients. Finally, caregivers reported a high increase in anxiety, depression, and distress.

**Conclusion:** Our study shows that quarantine for COVID-19 is associated with an acute worsening of clinical symptoms in patients with dementia as well as increase of caregivers' burden. Our findings emphasize the importance to implement new strategies to mitigate the effects of quarantine in patients with dementia.

**Keywords:** quarantine, COVID-19, dementia, Alzheimer's disease, BPSD, caregiver burden

## INTRODUCTION

The coronavirus disease 2019 (COVID-19) pandemic is, nowadays, a global public health emergency (Palacios Cruz et al., 2020). By December 8, 2020, over 66 million confirmed cases of COVID-19 with more than 1.530.000 deaths worldwide have been reported to the World Health Organization. Older adults and patients with certain comorbidities, many of whom are of advanced age, are particularly susceptible to more severe consequences of the disease (Shahid et al., 2020). The impact of the pandemic to the healthcare systems has been disruptive, and prevention as well as treatment services for patients with non-communicable diseases have been severely reduced.

Trying to slow down the spread of pandemic, several governments launched mitigation strategies, based mainly on quarantine. Quarantine is efficacious in reducing incidence and mortality during outbreaks of infectious diseases, and preliminary indications suggest that this strategy is effective also in the COVID-19 pandemic (Nussbaumer-Streit et al., 2020). However, home confinement is an unpleasant experience, due to the significant limitations in physical, cognitive, and social activities. Studies related to the 2003 outbreak of severe acute respiratory syndrome (SARS) in China and Canada, as well as the 2014 Ebola outbreak in Africa, showed that quarantine is associated with several negative psychological effects, like depression, irritability, anger, and insomnia (Hawryluck et al., 2004; Barbisch et al., 2015; Brooks et al., 2020; Gualano et al., 2020). Deaths by suicide increased in older adults during the SARS epidemic in Hong Kong (Cheung et al., 2008). In addition, long-lasting psychiatric effects of quarantine have been reported (Mak et al., 2009).

Individuals with Alzheimer's disease and other dementias are among the most vulnerable persons in the society, and the COVID-19 outbreak is likely to have further exacerbated their frailty (Wang et al., 2020). Quarantine effects in dementia patients and family caregivers have never been adequately investigated. All the negative effects of quarantine previously reported may be exacerbated in individuals with dementia, whose physical and social isolation may amplify the functional limitations and the pre-existing conflicts within the family. Patients with dementia

living at home significantly depend on family caregivers for assistance, and caring for dementia patients causes a significant burden in caregivers (Chiao et al., 2015; Raggi et al., 2015; Brown et al., 2020). Higher levels of cognitive, behavioral and motor impairment in patients with dementia are associated with greater burden and distress in their caregivers (Black et al., 2018; Yang et al., 2019; Xiong et al., 2020). Therefore, there is an urgent need to investigate clinical and psychological changes due to quarantine in patients with dementia and their caregivers.

This study aimed at evaluating the effects of quarantine in Italian patients with different types and severity of dementia and their caregivers. Approximately 45 days after quarantine declaration by the Italian government, we interviewed the family caregivers of persons with dementia referring to several Italian Centers for Cognitive Disorders and Dementia (CDCDs), investigating patients' variations in cognitive, behavioral, and motor symptoms. Besides, we evaluated the changes in prescribed medications, caregiver's burden, and changes in health services provided by the Italian National Health Service.

## MATERIALS AND METHODS

### Study Design, Centers and Participants

This is a multicentric, nation-wide survey. Eighty-nine Italian CDCDs were initially recruited for the study. First of all, a semi-structured questionnaire (see **Supplementary Material**) was administered to the Director of each Centre in order to evaluate its qualitative and quantitative characteristics, and variations in clinical activities after quarantine declaration. Due to the quarantine rules, all the out-patients visits were stopped. Therefore, the clinical staff members of each CDCD consecutively contacted by phone the family caregivers of patients registered in the waiting list. After an oral consent, a semi-structured, self-made interview gathering demographic and clinical data on the patient and the caregiver was administered (see **Supplementary Material**). Inclusion criteria were a diagnosis of one of the most common forms of dementias including: (1) Alzheimer's Disease (AD), (2) Dementia with Lewy Bodies (DLB), (3)

Frontotemporal Dementias (FTD), and (4) Vascular Dementia (VD). Exclusion criteria included current or previous diagnosis of other forms of dementias, mild cognitive impairment, and subjective cognitive complaints.

## Description of the Survey

We collected all the socio-demographic characteristics of patients. The stage of dementia was assessed using the Clinical Dementia Rating (CDR) (Morris et al., 1997) scale. Furthermore, we investigated the patient's variation in his/her clinical status, analyzing cognitive, psychological, behavioral, and motor symptoms during the quarantine period. Briefly, the following cognitive features were investigated: changes in memory, spatial and temporal orientation, language, attention, and perception. Questions about patient's awareness of the pandemic were administered. In addition, we acquired data regarding variations in behavioral and psychological symptoms of dementia (BPSD, i.e., irritability, apathy, agitation, anxiety, depression, sleep change, aggressiveness, wandering, appetite change, hallucinations, and delusions), reporting for these symptoms both a worsening and/or a new onset that occurred during the quarantine period. Further questions about the need for modifying therapy because of BPSD changes were administered. Finally, we investigated the changes in patients' motor activity, evaluating through a 5-point ordinal scale specific question if the subject walked better from the beginning of the quarantine period, remained stable, walked slower, became wheelchair, or became confined to bed.

## Caregivers' Socio-Demographic Characteristics

Caregivers' were asked about the cohabitation with the patient, composition of the family, and own work activity. Other questions concerned the impact of the quarantine period on caring for a person with dementia on their caregivers' lives, investigating both social and psychological effects (changes in caregiver's life, change in the relationship with the patient, concern for the pandemic, changes in therapeutic assistance, need to seek help from the emergency department, use of telemedicine, need to support during quarantine, own feeling of depression, anxiety, irritability, distress, overwhelm, and abandonment). Further details are available at **Supplementary Material**.

## Statistical Analysis

We performed statistical analysis using SPSS software, version 21, and the R statistical computing environment, version 3.6.2 (IBM Corp, 2012; R Core Team, 2013). Due to the low rate of missing data (<1%) no imputation was made. Firstly, we performed a descriptive analysis of all the demographic and clinical data. Then, we performed univariable and multivariable logistic regression of the dependent variables (outcomes in clinical symptoms, all described by binary variables) on the collected independent variables, using mixed effects logistic regression (as implemented in the lme4 R package) with the center as a random effect and all other regressors as fixed effects. Regressors with significant  $p$  ( $<0.05$ ) in univariable logistic regression were

included in multivariable regression. Bonferroni correction was applied to all the  $p$ -values of multivariable analysis, considering all outcomes together, as multivariable analysis is considered as our final result. We controlled for caregiver stress by including it in the multivariable analysis as a confounder, as a numerical fixed effect. The level of statistical significance was set at  $p < 0.05$ .

## Ethical Standards

The study was initially approved by the Ethics Committee of the Coordinating Centre (University of Torino on April 7, 2020, n.00150/2020) and then by the local ethics boards. Caregivers gave first oral and then written fully informed consent to the study.

## RESULTS

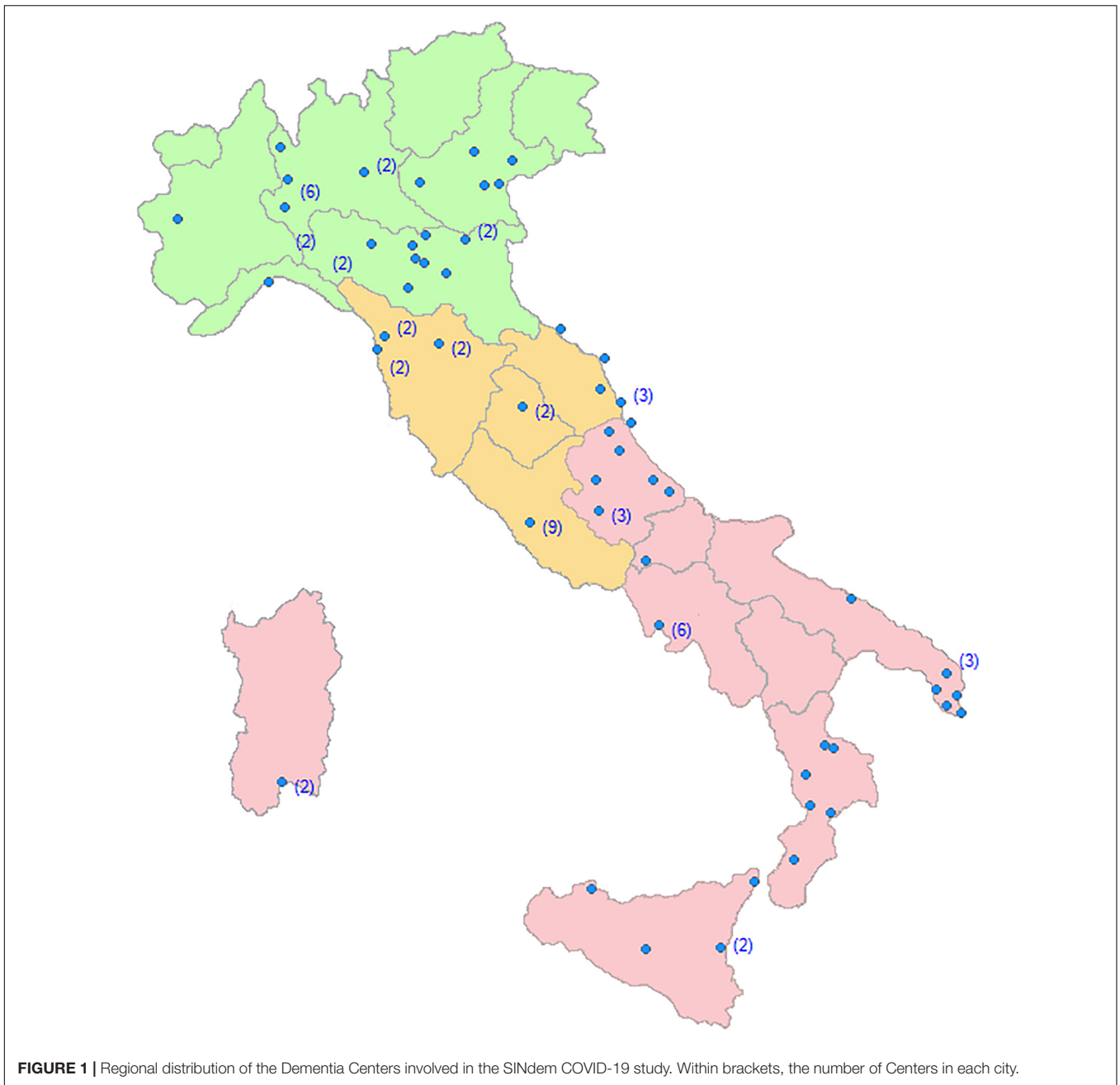
### Changes in Health Services

The response rate of CDCDs to the proposed questionnaires was 98%. Only two CDCDs were excluded because they did not recruit patients. The final 87 recruiting Centers were homogeneously distributed throughout the Italian territory (**Figure 1**).

Thirty-two percent of patients with dementia were recruited in Northern, 32% in Central, and 36% in Southern Italy. Thirty percent of CDCDs was based at university hospitals, 34% to general hospitals, and 36% to territorial based health services. In the pre-COVID-19 pandemic period, each Center followed a mean of 160 dementia patients per month. After the quarantine declaration, 85% of the CDCD suspended medical and psychological appointments, and visits were restricted to emergencies. Different forms of telemedicine (from phone calls to videoconferencing) were activated in 78% of Centers. One out of two Center provided on-line psychological support for caregivers. All the randomized clinical trials were stopped. Finally, 94% of support activities for dementia patients (Alzheimer's café, Day Center, etc.) were closed.

### Caregivers

The Survey response rates of the family caregivers ranged from 91.3 to 99.1%, according to different dementia Centers. We interviewed 5321 caregivers of patients regularly followed at different CDCDs. After data cleaning, the information collected from 4913 family caregivers (women 2934, mean age  $\pm$  SD = 58.2  $\pm$  12.0 years; men 1979, mean age  $\pm$  SD = 60.7  $\pm$  13.9 years) were analyzed. Fifty-nine % were cohabitants with the patients, and 36% were spouses of the dementia care recipient. Approximately half of the caregivers reported that quarantine induced a significant change in their lifestyle, with 30.3% complaining a reduction in time devoted to their own activities, and 15.5% reporting an increase in intrafamilial psychological conflicts. Caregivers reported a significant increase in anxiety (45.9%), depression (18.6%), irritability (26.2%), and distress (28.9%). Finally, 80.8% of the caregivers reported that telemedicine has been of help during quarantine.



## Dementia Patients

From April 14 to April 27, 2020, we collected data regarding 4913 patients with dementia (2934 females, mean age  $\pm$  SD = 78.9  $\pm$  8.2 years; 1979 males, mean age  $\pm$  SD = 77.2  $\pm$  8.0 years) regularly followed at CDCDs. Demographic and clinical characteristics of dementia patients are shown in **Table 1**.

Family caregivers reported that, after a quarantine period of approximately 47 days, dementia patients showed a worsening in cognitive symptoms (+55.1%), and behavioral symptoms (+51.9%), the onset of new behavioral symptoms (+25.9%), and an increase in motor symptoms (+36.7%). According to

caregivers, 40.1% of patients were totally aware of quarantine declaration, 36.0% were only partially aware, while 23.9% were totally unaware. Awareness of the quarantine varied significantly among the dementia subgroups, with DLB patients showing the highest degree (69.8%) of partial of total unawareness of pandemic.

## Changes in Cognitive Symptoms

Caregivers reported a worsening in cognitive symptoms in approximately 60% of dementia participants. About 37% of dementia participants showed worsening in two or more cognitive domains. Deficits more often reported as increased

**TABLE 1** | Demographic and clinical characteristics of study participants.

Patients	Total (n = 4913)	AD (n = 3372)	DLB (n = 360)	FTD (n = 415)	VD (n = 766)
Age (years, mean ± SD)	78.3 ± 8.2	78.3 ± 8.0	78 ± 7.3	72.3 ± 8.9	81.6 ± 7.0
Sex (female, %)	59.7	63.5	42.2	46.7	58.4
Duration of the disease (years, mean ± SD)	4.5 ± 3.1	4.6 ± 3.1	4.5 ± 3	4.8 ± 3.2	4.1 ± 2.9
<b>Regional distribution</b>					
North (%)	32.2	26.5	35.3	47.5	47.8
Centre (%)	31.5	34.1	36.4	21.2	23.4
South - Islands (%)	36.3	39.4	28.3	31.3	28.8
<b>CDR Stage (%)</b>					
1	25.0	24.3	26.3	23.4	28.4
2	47.8	49.2	41.9	48.6	43.8
3	27.2	26.5	31.8	28.0	27.8
Duration of the quarantine (days, mean ± SD)	47.2 ± 6.4	47.2 ± 6.5	46.6 ± 5.6	46.4 ± 5.3	47.8 ± 6.8
<b>Changes in cognitive symptoms (Yes, %)</b>					
Sex (female, %)	58.4	63.0	36.3	44.7	55.2
<b>Changes in BPSD (Yes, %)</b>					
Sex (female, %)	57.9	62.9	38.4	45.4	55.1
<b>New onset BPSD (Yes, %)</b>					
Sex (female, %)	56.7	59.8	41.7	41.8	56.1
<b>Changes in motor symptoms (Yes, %)</b>					
Sex (female, %)	58.6	65.0	44.7	43.1	52.5
<b>Caregivers</b>					
Age (years, mean ± SD)	59.3 ± 13	59.3 ± 13.1	60.7 ± 12.7	59.1 ± 13.6	60 ± 12.4
Sex (female, %)	53.9	51.2	66.4	55.4	59.4
Cohabitant caregiver (%)	58.9	58.1	63.5	69.6	54.4
<b>Degree of kinship</b>					
Spouses (%)	36.0	35.0	43.1	54.8	26.9
Son/daughter (%)	54.5	55.5	48.7	37	62.5
Others (%)	9.5	9.5	8.2	8.2	10.6
Increase in anxiety (%)	45.9	46.1	43.4	44.2	47.4
Increase in depression (%)	18.6	17.2	21.3	24.3	20.3

AD, Alzheimer's disease; DLB, Dementia with Lewy Bodies; FTD, frontotemporal dementia; VD, vascular dementia; CDR, Clinical Dementia Rating; BPSD, Behavioral and Psychological symptoms of dementia.

were: forgetfulness (68%), confusion (67.9%), and temporal disorientation (37%). **Figure 2** displays the frequency distribution of cognitive symptoms worsened during quarantine.

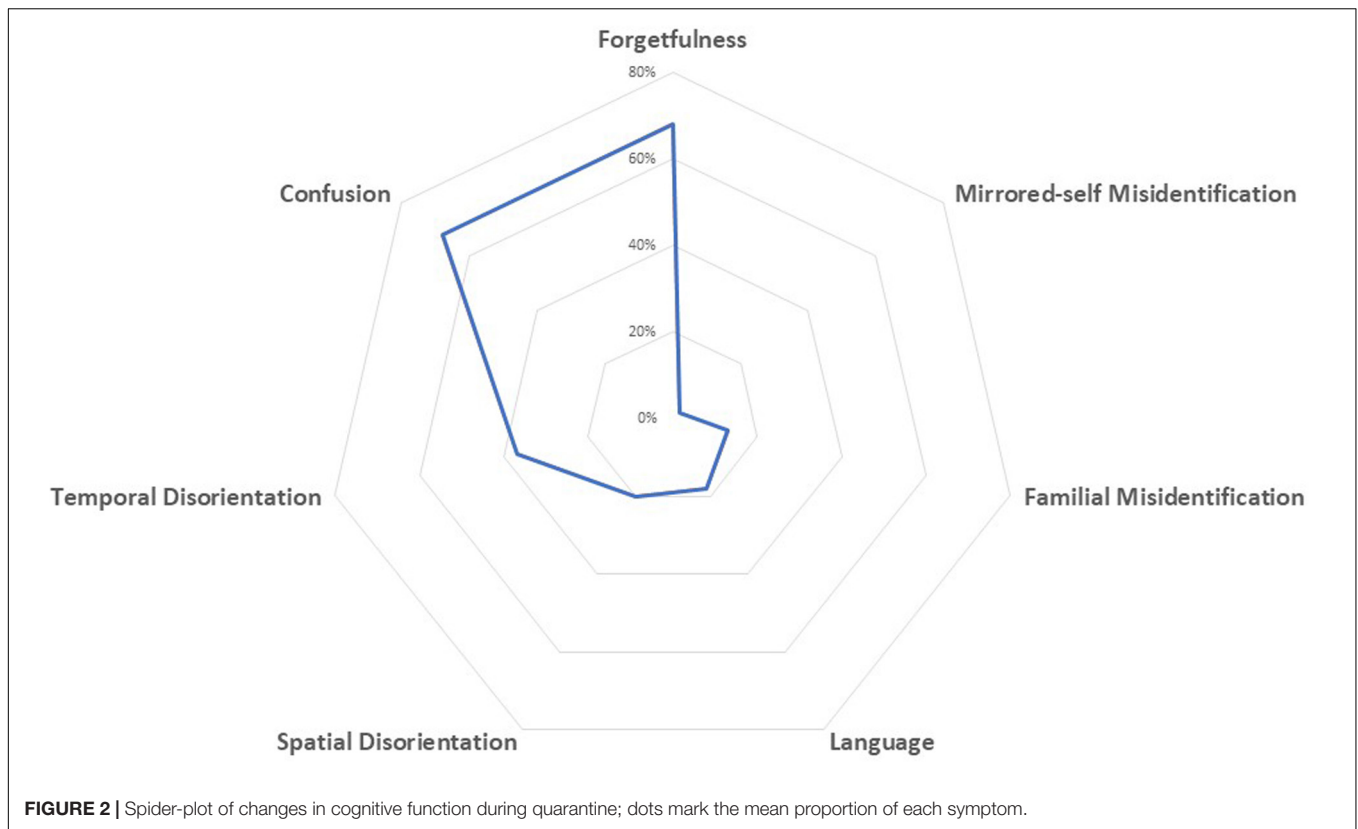
When we examined diagnostic subgroups, we found a worsening of cognitive symptoms mainly in patients with DLB (59.6%) and AD (55.7%), followed by VD (54.2%), and FTD (48.3%) patients. Thirty-eight % of patients with AD showed a worsening in memory functions, 37.5% presented an increase in confusion, and 21.1% showed a worsening in temporal disorientation. The same effects were observed in patients with DLB, showing a worsening in memory functions in 26.1%, an increase in confusion in 40.3%, and in temporal disorientation in 20.3%. Patients with FTD showed mainly a worsening in language functions (25.5%). After multivariable analysis, the increase of cognitive symptoms in the overall group of patients with dementia was not associated with age, gender, duration of quarantine, disease severity, and changes in caregivers' psychological features. Contrariwise, an inverse relationship between the aggravation of cognitive symptoms and the duration of the dementia was found ( $p < 0.001$ ).

Interestingly, the pre-existing total or partial independence in motor function represents a risk factor for the worsening of cognitive functions (OR 1.85 [95% CI 1.42–2.39],  $p < 0.001$ ; and OR 2.08 [95% CI 1.68–2.57],  $p < 0.001$ , respectively). Contrariwise, total awareness of quarantine was a protective factor against cognitive worsening (OR 0.68 [95% CI 0.57–0.81],  $p < 0.001$ ) (**Table 2**).

## Changes in Behavioral Symptoms

During the quarantine period, BPSD worsened in approximately half of the patients with dementia. The most frequently worsened BPSD were irritability (40%), followed by apathy (35%), and agitation (31%) (**Figure 3**).

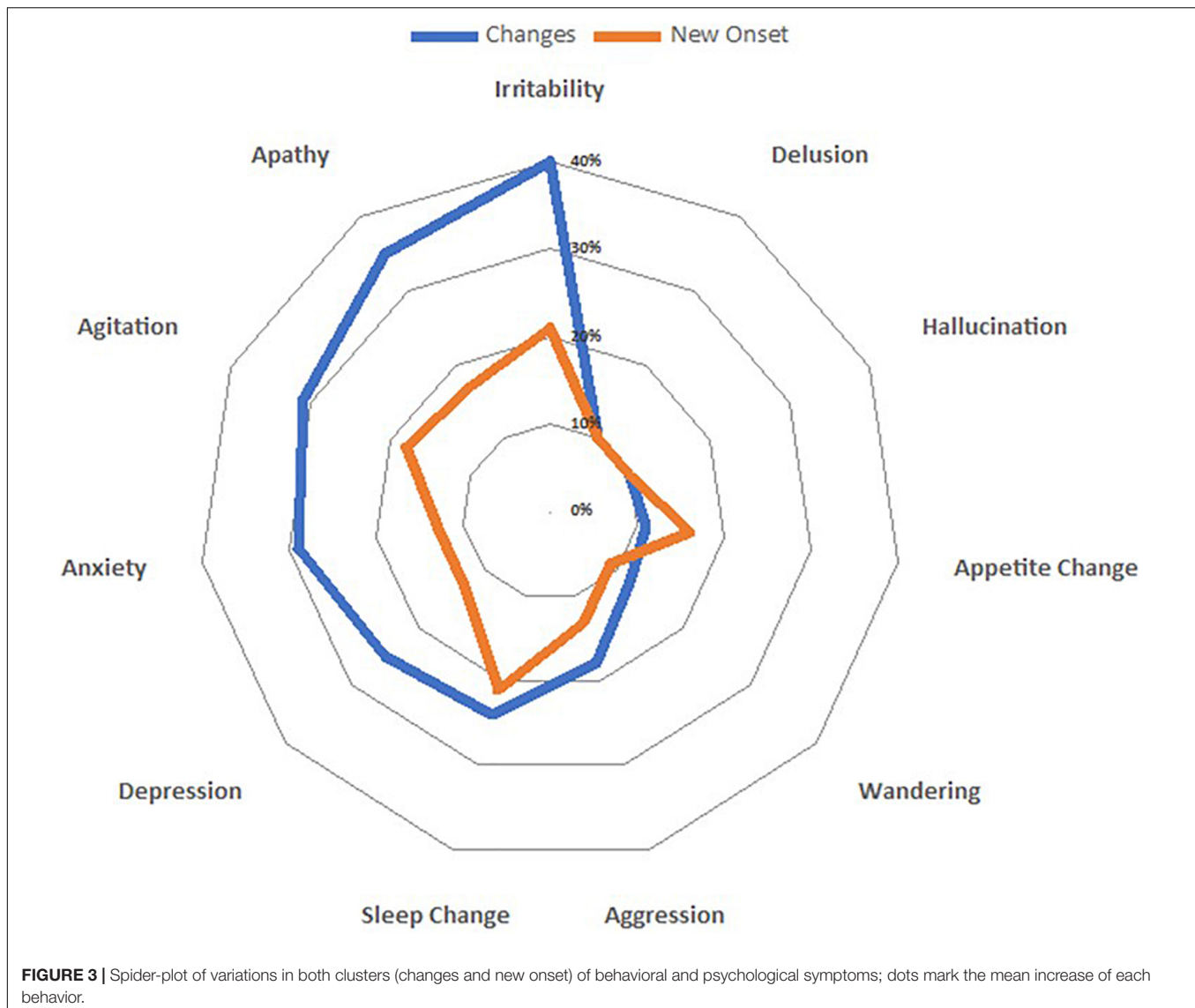
Examining subgroups, we found an increase in the number of symptoms mainly in patients with DLB (63.8%), followed by FTD (55.3%), AD (50.5%), and VD (50.3%). The increase of neuropsychiatric symptoms in patients with dementia was associated with moderate disease severity (CDR 2,  $p = 0.009$ ), but not remained significant after applying Bonferroni correction. The worsening of BPSD was associated with pre-existing total



**TABLE 2 |** Univariable and multivariable analysis of risk and protective factors for cognitive and motor changes during quarantine in patients with dementia.

	Univariable analysis		Multivariable analysis		P-value after Bonferroni correction
	Crude OR (95% CI)	P-value	Adjusted OR (95% CI)	P-value	
<b>Changes in cognitive symptoms</b>					
Age	1.00 (0.99 – 1.00)	0.24			
Sex (men vs. women)	1.15 (1.01 – 1.30)	0.03	1.13 (0.99 – 1.29)	0.07	1
Duration of the disease (years)	0.95 (0.93 – 0.97)	<0.001	0.94 (0.92 – 0.97)	<0.001	<0.001
Duration of quarantine (days)	1.00 (0.99 – 1.01)	0.97			
Disease severity – CDR 2	1.29 (1.11 – 1.50)	<0.001	1.19 (1.00 – 1.42)	0.05	1
Disease severity – CDR 3	1.02 (0.86 – 1.21)	0.81	0.93 (0.74 – 1.17)	0.52	1
Total physical independence	1.74 (1.40 – 2.16)	<0.001	1.85 (1.42 – 2.39)	<0.001	<0.001
Partial physical independence	2.19 (1.80 – 2.67)	<0.001	2.08 (1.68 – 2.57)	<0.001	<0.001
Total awareness of pandemic	0.74 (0.65 – 0.85)	<0.001	0.68 (0.57 – 0.81)	<0.001	<0.001
Partial awareness of pandemic	1.39 (1.18 – 1.64)	<0.001	1.20 (1.00 – 1.43)	0.05	1
<b>Changes in motor symptoms</b>					
Age	1.03 (1.02 – 1.04)	<0.001	1.02 (1.01 – 1.03)	<0.001	0.001
Sex (men vs. women)	1.11 (0.98 – 1.25)	0.12			
Duration of the disease (years)	1.05 (1.03 – 1.07)	<0.001	1.00 (0.98 – 1.03)	0.86	1
Duration of quarantine (days)	1.00 (0.99 – 1.01)	0.99			
Disease severity – CDR 2	1.94 (1.65 – 2.29)	<0.001	1.29 (1.07 – 1.56)	0.009	0.29
Disease severity – CDR 3	3.08 (2.56 – 3.70)	<0.001	1.60 (1.26 – 2.02)	<0.001	0.004
Total physical independence	0.32 (0.25 – 0.40)	<0.001	0.49 (0.37 – 0.64)	<0.001	<0.001
Partial physical independence	0.93 (0.77 – 1.12)	0.44	1.06 (0.86 – 1.30)	0.59	1
Total awareness of pandemic	0.44 (0.38 – 0.51)	<0.001	0.76 (0.64 – 0.91)	0.003	0.10
Partial awareness of pandemic	0.75 (0.64 – 0.88)	<0.001	0.93 (0.78 – 1.10)	0.39	1

OR, Odds ratio; 95% CI, 95% confidence interval; CDR, Clinical Dementia Rating.



or partial autonomy in motor function (totally independent OR 1.84 [95% CI 1.43–2.38],  $p < 0.001$ ; partially dependent OR 1.88 [95% CI 1.53–2.31],  $p < 0.001$ ). After multivariate analysis, the worsening of BPSD was not associated with age, gender, duration of quarantine, disease severity, duration of disease, and changes in caregivers' psychological features. On the contrary, total awareness of the pandemic was inversely related to worsening of BPSD (OR 0.75 [95% CI 0.63–0.89],  $p = 0.03$ ) (Table 3).

New BPSD occurred in approximately a quarter of the dementia patients. There was a different increase in the frequency of BPSD across groups. The most frequently reported new symptom was irritability (21.3%), followed by sleep change (21%), and agitation (18%) (Figure 2), which required therapy adjustments. The highest increase was observed in AD (26.7%), intermediate in VD and DLB (25.6 and 23.3%, respectively), and the lowest increase in FTD (21.9%). No suicide attempts were reported. The onset of new BPSD was associated with the total or partial physical independence ( $p < 0.001$  and  $p = 0.01$ ,

respectively). Total awareness of the pandemic was inversely related to onset of new BPSD (OR 0.74, [95% CI 0.62–0.88],  $p = 0.03$ ).

### Changes in Motor Symptoms

Increased motor dysfunction, mainly characterized by walking difficulties, was reported by caregivers in 36.7% of the patients. There was an increase mainly in patients with DLB (52.8%), followed by VD (42.3%), FTD (40.2%), and AD (33.2%). Age and disease severity were associated with an increase of motor dysfunction ( $p < 0.001$ ) during quarantine, whereas full physical autonomy before quarantine was a positive predictor against worsening in motor functions (OR 0.49 [95% CI 0.37–0.64],  $p < 0.001$ ).

The increased prevalence of cognitive, behavioral and motor symptoms was not influenced either by regional distribution of different CDCD or by the regional prevalence of COVID-19 at the time of the study.

**TABLE 3 |** Univariable and multivariable analysis of risk and protective factors for increase in BPSD and onset of new BPSD during quarantine in patients with dementia.

	Univariable analysis		Multivariable analysis		
	Crude OR (95% CI)	P-value	Adjusted OR (95% CI)	P-value	P-value after Bonferroni correction
<b>Increase in BPSD</b>					
Age	0.99 (0.98 – 0.99)	<0.001	0.99 (0.98 – 0.99)	0.005	0.16
Sex (men vs. women)	1.17 (1.04 – 1.32)	0.008	1.12 (0.99 – 1.28)	0.08	1
Duration of the disease (years)	0.98 (0.96 – 0.99)	0.04	0.97 (0.95 – 0.99)	0.01	0.48
Duration of quarantine (days)	0.99 (0.99 – 1.01)	0.66			
Disease severity – CDR 2	1.33 (1.15 – 1.54)	<0.001	1.26 (1.06 – 1.49)	0.009	0.30
Disease severity – CDR 3	1.32 (1.12 – 1.56)	0.001	1.21 (0.97 – 1.52)	0.09	1
Total physical independence	1.56 (1.26 – 1.93)	<0.001	1.84 (1.43 – 2.38)	<0.001	<0.001
Partial physical independence	1.90 (1.57 – 2.30)	<0.001	1.88 (1.53 – 2.31)	<0.001	<0.001
Total awareness of pandemic	0.72 (0.63 – 0.82)	<0.001	0.75 (0.63 – 0.89)	<0.001	0.03
Partial awareness of pandemic	1.25 (1.07 – 1.45)	0.005	1.20 (1.01 – 1.42)	0.04	1
<b>Onset of new BPSD</b>					
Age	1.00 (0.99 – 1.00)	0.32			
Sex (men vs. women)	1.19 (1.04 – 1.36)	0.01	1.10 (0.95 – 1.27)	0.21	1
Duration of the disease (years)	0.99 (0.97 – 1.01)	0.41			
Duration of quarantine (days)	1.00 (0.99 – 1.01)	0.72			
Disease severity – CDR 2	1.14 (0.96 – 1.35)	0.13			
Disease severity – CDR 3	1.08 (0.89 – 1.32)	0.41			
Total physical independence	1.69 (1.30 – 2.18)	<0.001	1.92 (1.45 – 2.55)	<0.001	<0.001
Partial physical independence	1.59 (1.26 – 2.02)	<0.001	1.55 (1.22 – 1.99)	<0.001	0.01
Total awareness of pandemic	0.78 (0.67 – 0.92)	0.003	0.74 (0.62 – 0.88)	<0.001	0.03
Partial awareness of pandemic	1.17 (0.99 – 1.39)	0.07	1.10 (0.92 – 1.32)	0.29	1

BPSD, Behavioral and Psychological Symptoms of Dementia; OR, Odds ratio; 95% CI, 95% confidence interval; CDR, Clinical Dementia Rating.

## DISCUSSION

Our study shows that quarantine due to COVID-19 pandemic is associated with a dramatic increase in clinical symptoms of patients with dementia. According to family caregivers, social isolation and physical restraint caused a worsening in cognitive function, an aggravation of several behavioral symptoms, and a worsening in motor function. These effects were observed in all forms of dementia, but patients with DLB and AD showed the highest increase in cognitive and behavioral symptoms. In addition, the quarantine period was associated with an increase of caregiver's burden, mainly characterized by anxiety.

To the best of our knowledge, this is the first nationwide study that investigated the effects of quarantine in patients with dementia. Our data confirm the results of a previous, small study that investigated the effects of COVID-19 outbreak in patients with MCI and dementia (Canevelli et al., 2020). To evaluate factors associated with the quarantine-related clinical worsening, we performed extensive statistical analyses, and the most consistent relation was observed with the pre-pandemic physical status. Patients still able to walk unassisted or accompanied, after an acute interruption of physical activity, showed the greater worsening in clinical symptoms. Sex, age, duration of the disease, and disease severity had small and inconsistent effects. Several studies showed that physical activity has a positive impact on cognition of elderly adults (Fratiglioni et al., 2004; Bherer, 2015; Forbes et al., 2015; Young et al., 2015) and provided evidence that

in patients with dementia there is an inverse relationship between physical activity, abnormal behavior, and cognitive decline (Nelson and Tabet, 2015; Karssemeijer et al., 2017). Patients physically more independent and, probably, with a more active life, enriched with a variety of social contacts and cognitive stimulation, are those most affected by the negative effects of confinement. Our study highlights the importance of maintaining physical activity in patients with dementia, and suggests the need for additional investigations to better understand this phenomenon.

Our study showed that 40% of the examined patients were totally aware of the COVID-19 pandemic and of social and physical restrictive measures planned to control virus transmission. Intriguingly, we found that patients with dementia still able to completely understand the need of quarantine, and therefore able to adopt adequate strategies to face the stress, had a significantly lower impairment in cognitive, behavioral and motor function. Contrariwise, less aware patients failed in creating new adaptive procedures to quarantine. Awareness of quarantine varied significantly among dementia subgroups and DLB patients showed the highest degree of partial and total degree of awareness. Taken together, the results of our study suggested that DLB patients are the most vulnerable to acute stress among the patients with dementia.

The neurobiological basis of the observed phenomenon has never been investigated. Examining subgroups of patients with dementia, we found an increase mainly of the core symptoms that characterize different types of dementing illness. This finding



suggests that the observed phenomenon is not disease-specific but is more likely attributable to stress-induced changes. Studies in experimental animals showed that exposure to both acute and chronic stress evokes chemical changes in brain that impair the higher cognitive functions (Arnsten et al., 1999; Yamada et al., 2003). Uncontrollable, acute stress induces a diffuse increase in brain glucocorticoids and a specific catecholamine release in prefrontal cortex, impairing spatial memory tasks (Arnsten, 2015; Bahtiyar et al., 2020). Intriguingly, in animal models of acute physical and social restraints, an impairment of serotonin metabolism in the central nervous system was reported, with an alteration of immune response (Medina-Martel et al., 2013). In humans, stress drives several diseases, including cognitive disorders such as AD (Canet et al., 2019). Acute stress induces a neuroinflammatory response characterized by the release of several pro-inflammatory molecules and microglial activation that is a common feature of several neurodegenerative diseases (Sharma and Kanneganti, 2016). Finally, worsening of sleep, frequently reported in our study, is an additional explanation of our findings. Poor sleep is strongly associated with risk of multiple types of dementia and, via modulation of  $\beta$ -amyloid secretion, is directly involved in AD pathogenesis (Vanderheyden et al., 2018). Further studies are warranted to better elucidate the neurobiological mechanisms underlying the reported worsening of patients with dementia during quarantine and to provide therapeutic strategies.

Caregiving of dementia patients is associated with a higher prevalence of depressive and anxiety disorders, and impairments in physical health. Data from our study clearly showed that, as previously observed in the general population after SARS and Ebola outbreaks, quarantine is associated with increase in anxiety and depression. New health policies should be planned in the post-COVID-19 era to help family caregivers in delivering effective care. Finally, dementia Centers need to implement new strategies, as telemedicine (Goodman-Casanova et al., 2020; Hollander and Carr, 2020), in order to assist frailty people and support family members.

Some limitations of our study should be acknowledged. We examined dementia patients living at home and our data cannot be generalized to institutionalized patients with dementia. Then, we performed a cross-sectional study and, at present, it is unclear whether the observed clinical worsening is a transient or long-lasting phenomenon. Finally, all the clinical data regarding patient's symptoms were collected from family caregivers as, due to quarantine rules, it was not possible to administer face-to-face standardized neuropsychological tests. However, previous studies showed that BPSD can be adequately evaluated by family caregivers and demonstrated that symptoms reported can be used in the clinical staging of dementia (Kwok et al., 2011; Yuan et al., 2020). Therefore, in order to evaluate if there is a time-effect, we have planned a new follow-up interview to the same caregivers. Finally, the major strength of our study is the high number of caregivers that have been interviewed in a relatively small amount of time and their reliability, having this role from the onset of dementia.

## CONCLUSION

In conclusion, interviewing family caregivers, we investigated for the first time the acute effects of quarantine in a large Italian population of patients with dementia, and we found that restrictive public health measures, are associated with an increase of dementia symptoms. These findings are important from a health policy perspective and suggest the need of new health care strategies to target patients with dementia during pandemic.

## DATA AVAILABILITY STATEMENT

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

## ETHICS STATEMENT

The studies involving human participants were reviewed and approved by Comitato Etico Interaziendale di Torino. The patients/participants provided their written informed consent to participate in this study.

## AUTHOR CONTRIBUTIONS

IR, AB, CM, AC, LB, CC, and VL designed the study and planned center recruitment. AC, CM, ER, and AB wrote the report. RDL and PP did the statistical analyses. ER, AV, VI, NV, FA, IA, PC, CB, RS, DQ, VG, GL, MF, GT, and CF contributed to the interpretation and discussion of results and reviewed the manuscript. The collaborating authors contributed to the collection of clinical data. All the authors and the collaborating authors contributed to the article and approved the submitted version.

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## SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fnagi.2020.625781/full#supplementary-material>

## REFERENCES

- Arnsten, A. F. (2015). Stress weakens prefrontal networks: molecular insult to higher cognition. *Nat. Neurosci.* 18, 1376–1385. doi: 10.1038/nn.4087
- Arnsten, A. F. T., Mathew, R., Ubriani, R., Taylor, J. R., and Li, B. M. (1999). Alpha-1 noradrenergic receptor stimulation impairs prefrontal cortical cognitive function. *Biol. Psychiatry* 45, 26–31. doi: 10.1016/s0006-3223(98)00296-0
- Bahtiyar, S., Gulmez Karaca, K., Henckens, M. J. A. G., and Roozendaal, B. (2020). Norepinephrine and glucocorticoid effects on the brain mechanisms underlying memory accuracy and generalization. *Mol. Cell. Neurosci.* 108:103537. doi: 10.1016/j.mcn.2020.103537
- Barbisch, D., Koenig, K. L., and Shih, F. Y. (2015). Is there a case for quarantine? Perspectives from SARS to Ebola. *Disaster Med. Public Health Prep.* 9, 547–553. doi: 10.1017/dmp.2015.38
- Bherer, L. (2015). Cognitive plasticity in older adults: effects of cognitive training and physical exercise. *Ann. N.Y. Acad. Sci.* 1337, 1–6. doi: 10.1111/nyas.12682
- Black, C. M., Ritchie, C. W., Khandker, R. K., Nalder, E., Cameron, J. I., Mihailidis, A., et al. (2018). Non-professional caregiver burden is associated with the severity of patients' cognitive impairment. *PLoS One* 13:e0204110. doi: 10.1371/journal.pone.0204110
- Brooks, S. K., Webster, R. K., Smith, L. E., Woodland, L., Wessely, S., Greenberg, N., et al. (2020). The psychological impact of quarantine and how to reduce it: rapid review of the evidence. *Lancet* 395, 912–920. doi: 10.1016/S0140-6736(20)30460-8
- Brown, E. E., Kumar, S., Rajji, T. K., Pollock, B. G., and Mulsant, B. H. (2020). Anticipating and mitigating the impact of the COVID-19 pandemic on Alzheimer's disease and related dementias. *Am. J. Geriatr. Psychiatry* 28, 712–721. doi: 10.1016/j.jagp.2020.04.010
- Canet, G., Hernandez, C., Zussy, C., Chevaller, N., Desrumaux, C., and Givalois, L. (2019). Is AD a stress-related disorder? Focus on the HPA axis and its promising therapeutic targets. *Front. Aging Neurosci.* 11:269. doi: 10.3389/fnagi.2019.00269
- Canevelli, M., Valletta, M., Toccaceli Blasi, M., Remoli, G., Sarti, G., Nuti, F., et al. (2020). Facing dementia during the COVID-19 outbreak. *J. Am. Geriatr. Soc.* 68, 1673–1676. doi: 10.1111/jgs.16644
- Cheung, Y. T., Chau, P. H., and Yip, P. (2008). A revisit on older adults suicides and Severe Acute Respiratory Syndrome (SARS) epidemic in Hong Kong. *Int. J. Geriatr. Psychiatry* 23, 1231–1238. doi: 10.1002/gps.2056
- Chiao, C. Y., Wu, H. S., and Hsiao, C. Y. (2015). Caregiver burden for informal caregivers of patients with dementia: a systematic review. *Int. Nurs. Rev.* 62, 340–350. doi: 10.1111/inr.12194
- Forbes, D., Forbes, S. C., Blake, C. M., Thiessen, E. J., and Forbes, S. (2015). Exercise programs for people with dementia. *Cochrane Database Syst. Rev.* 4:CD006489. doi: 10.1002/14651858.CD006489
- Fratiglioni, L., Paillard-Borg, S., and Winblad, B. (2004). An active and socially integrated lifestyle in late life might protect against dementia. *Lancet Neurol.* 3, 343–353. doi: 10.1016/S1474-4422(04)00767-7
- Goodman-Casanova, J. M., Dura-Perez, E., Guzman-Parra, J., Cuesta-Vargas, A., and Mayoral-Cleries, F. (2020). Telehealth home support during COVID-19 confinement for community-dwelling older adults with mild cognitive impairment or mild dementia: survey study. *J. Med. Internet. Res.* 22:e19434. doi: 10.2196/19434
- Gualano, M. R., Lo Moro, G., Voglino, G., Bert, F., and Siliquini, R. (2020). Effects of Covid-19 lockdown on mental health and sleep disturbances in Italy. *Int. J. Environ. Res. Public Health* 17:E4779. doi: 10.3390/ijerph17134779
- Hawryluck, L., Gold, W. L., Robinson, S., Pogorski, S., Galea, S., and Styr, R. (2004). SARS control and psychological effects of quarantine, Toronto, Canada. *Emerg. Infect. Dis.* 10, 1206–1212. doi: 10.3201/eid1007.030703
- Hollander, J. E., and Carr, B. G. (2020). Virtually Perfect? Telemedicine for Covid-19. *N. Engl. J. Med.* 382, 1679–1681. doi: 10.1056/NEJMp2003539
- IBM Corp (2012). *Released 2012. IBM SPSS Statistics for Windows, Version 21.0.* Armonk, NY: IBM Corp.
- Karssemeijer, E. G. A., Aaronson, J. A., Bossers, W. J., Smits, T., Olde Rikkert, M. G. M., and Kessels, R. P. C. (2017). Positive effects of combined cognitive and physical exercise training on cognitive function in older adults with mild cognitive impairment or dementia: a meta-analysis. *Ageing Res. Rev.* 40, 75–83. doi: 10.1016/j.arr.2017.09.003
- Kwok, Y. T., Chen, C. Y., Chiu, M. J., Tang, L. Y., and Leung, K. K. (2011). Assessment of behavioural and psychological symptoms of dementia by family caregivers. *Arch. Gerontol. Geriatr.* 52, 60–65. doi: 10.1016/j.archger.2010.01.021
- Mak, I. W., Chu, C. M., Pan, P. C., Yiu, M. G., and Chan, V. L. (2009). Long-term psychiatric morbidities among SARS survivors. *Gen. Hosp. Psychiatry* 31, 318–326. doi: 10.1016/j.genhosppsych.2009.03.001
- Medina-Martel, M., Urbina, M., Fazzino, F., and Lima, L. (2013). Serotonin transporter in lymphocytes of rats exposed to physical restraint stress. *Neuroimmunomodulation* 20, 361–367. doi: 10.1159/000353797
- Morris, J. C., Ernesto, C., Schafer, K., Coats, M., Leon, S., Sano, M., et al. (1997). Clinical dementia rating training and reliability in multicentre studies: the Alzheimer's Disease Cooperative Study experience. *Neurology* 48, 1508–1510. doi: 10.1212/wnl.48.6.1508
- Nelson, L., and Tabet, N. (2015). Slowing the progression of Alzheimer's disease; what works? *Ageing Res. Rev.* 23, 193–209. doi: 10.1016/j.arr.2015.07.002
- Nussbaumer-Streit, B., Mayr, V., Dobrescu, A. I., Chapman, A., Persad, E., Klerings, I., et al. (2020). Quarantine alone or in combination with other public health measures to control COVID-19: a rapid review. *Cochrane Database Syst. Rev.* 4:CD013574. doi: 10.1002/14651858.CD013574
- Palacios Cruz, M., Santos, E., Velázquez Cervantes, M. A., and León Juárez, M. (2020). COVID-19, a worldwide public health emergency. *Rev. Clin. Esp.* doi: 10.1016/j.rce.2020.03.001 [Epub ahead of print].
- R Core Team (2013). *R: A Language and Environment for Statistical Computing.* Vienna: R Foundation for Statistical Computing.
- Raggi, A., Tasca, D., Panerai, S., Neri, W., and Ferri, R. (2015). The burden of distress and related coping processes in family caregivers of patients with Alzheimer's disease living in the community. *J. Neurol. Sci.* 358, 77–81. doi: 10.1016/j.jns.2015.08.024
- Shahid, Z., Kalayanamitra, R., McClafferty, B., Kepko, D., Ramgobin, D., Patel, R., et al. (2020). COVID-19 and older adults: What we know. *J. Am. Geriatr. Soc.* 68, 926–929. doi: 10.1111/jgs.16472
- Sharma, D., and Kanneganti, T. D. (2016). The cell biology of inflammasomes: mechanisms of inflammasome activation and regulation. *J. Cell Biol.* 213, 617–629. doi: 10.1083/jcb.201602089
- Vanderheyden, W. M., Lim, M. M., Musiek, E. S., and Gerstner, J. R. (2018). Alzheimer's disease and sleep-wake disturbances: amyloid, astrocytes, and animal models. *J. Neurosci.* 38, 2901–2910. doi: 10.1523/JNEUROSCI.1135-17.2017
- Wang, H., Li, T., Barbarino, P., Brodaty, H., Molinuevo, J. L., Xie, H., et al. (2020). Dementia care during COVID-19. *Lancet* 395, 1190–1191. doi: 10.1016/S0140-6736(20)30755-8
- Xiong, C., Biscardi, M., Astell, A., Nalder, E., Cameron, J. I., Mihailidis, A., et al. (2020). Sex and gender differences in caregiving burden experienced by family caregivers of persons with dementia: a systematic review. *PLoS One* 15:e0231848. doi: 10.1371/journal.pone.0231848
- Yamada, K., McEwen, B. S., and Pavlides, C. (2003). Site and time dependent effects of acute stress on hippocampal long-term potentiation in freely behaving rats. *Exp. Brain Res.* 152, 52–59. doi: 10.1007/s00221-003-1519-0

- Yang, F., Ran, M., and Luo, W. (2019). Depression of persons with dementia and family caregiver burden: finding positives in caregiving as a moderator. *Geriatr. Gerontol. Int.* 19, 414–418. doi: 10.1111/ggi.13632
- Young, J., Angevaren, M., Rusted, J., and Tabet, N. (2015). Aerobic exercise to improve cognitive function in older people without known cognitive impairment. *Cochrane Database Syst. Rev.* 4:CD005381. doi: 10.1002/14651858.CD005381
- Yuan, Q., Tan, T. H., Wang, P., Devi, F., Ong, H. L., Abidin, E., et al. (2020). Staging dementia based on caregiver reported patient symptoms: implications from a latent class analysis. *PLoS One* 15:e0227857. doi: 10.1371/journal.pone.0227857

**Conflict of Interest:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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