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## Associations of subjective and objective cognitive functioning after COVID-19: A six-month follow-up of ICU, ward, and home-isolated patients

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### ABSTRACT

**Background:** Subjective and objective cognitive dysfunction are reported after COVID-19 but with limited data on their congruence and associations with the severity of the acute disease. The aim of this cohort study is to describe the prevalence of subjective and objective cognitive dysfunction at three and six months after COVID-19 and the associations of subjective cognitive symptoms and psychological and disease-related factors.

**Methods:** We assessed a cohort of 184 patients at three and six months after COVID-19: 82 patients admitted to the Intensive Care Unit (ICU), 53 admitted to regular hospital wards, and 49 isolated at home. A non-COVID control group of 53 individuals was included. Demographic and clinical data were collected. Subjective cognitive symptoms, objective cognitive impairment, and depressive and post-traumatic stress disorder (PTSD) symptoms were assessed.

**Results:** At six months, subjective cognitive impairment was reported by 32.3% of ICU-treated, 37.3% of ward-treated, and 33.3% of home-isolated patients and objective cognitive impairment was observed in 36.1% of ICU-treated, 34.7% of ward-treated, and 8.9% of home-isolated patients. Subjective cognitive symptoms were associated with depressive and PTSD symptoms and female sex, but not with objective cognitive assessment or hospital metrics.

**Conclusions:** One-third of COVID-19 patients, regardless of the acute disease severity, reported high levels of subjective cognitive dysfunction which was not associated with results from objective cognitive screening but with psychological and demographic factors. Our study stresses the importance of thorough assessment of patients reporting long-term subjective symptoms, screening for underlying mental health related factors such as PTSD or depression.

### 1. Introduction

COVID-19 can have sequelae lasting for months after initial recovery (Lopez-Leon et al., 2021), including neurological (Chou et al., 2021) and psychiatric (Premraj et al., 2022) manifestations and cognitive impairment (Daroische et al., 2021). Cognitive impairment is one of the most common prolonged symptoms of COVID-19, as defined by the WHO consensus on post COVID-19 condition (WHO, 2021; Davis et al., 2021). The incidence of cognitive symptoms has ranged from 0 to 78% (Schou et al., 2021) in studies of the acute and post-acute phases of COVID-19

(Krishnan et al., 2022; Zhou et al., 2020). The heterogeneity of study samples and methods may partly explain the large variability in the results. Moreover, few studies have included a control group (Tavar-es-Junior et al., 2022), which would facilitate differentiating direct effects of the disease and indirect factors of the pandemic (Schou et al., 2021).

There is still limited information on the long-term cognitive outcome of patients with different severity levels of the acute disease (Hampshire et al., 2021), and particularly in mild COVID-19 (Schou et al., 2021; Van Kessel et al., 2022; Vanderlind et al., 2021). Patients who have been

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treated in the intensive care unit (ICU) have been found to have more cognitive impairment compared to other hospitalized patients (Taquet et al., 2021), although no differences between these patient groups have been found in a three-month follow-up (Rass et al., 2022). Compared to home-treated patients, hospitalization may be a risk to having more cognitive impairments (Becker et al., 2021). However, persisting cognitive symptoms and mental distress may appear also among patients with mild COVID-19 (Van Kessel et al., 2022).

Cognitive functioning can be evaluated using objective assessment or subjective self-reporting questionnaires. COVID-19 patients have commonly reported subjective cognitive complaints (Vanderlind et al., 2021) with as much as 54.7% reported in a 1-year follow-up (Ferrucci et al., 2022). However, there are contradictory results on how self-reported cognitive issues reflect objective cognitive evaluation (Almeria et al., 2020; Blackmon et al., 2022; Miskowiak et al., 2021). Besides infection-related factors, other mechanisms such as education, age, and psychological factors including distress may modulate cognitive outcomes (Majoka and Schimming, 2021). COVID-19 survivors may experience mental distress, including symptoms of depression (Renaud-Charest et al., 2021) and post-traumatic stress disorder (PTSD) (Heesakkers et al., 2022). Because of the symptomatic heterogeneity of COVID-19, studies employing a holistic approach combining clinical assessments with patient-reported measures and medical data are called for (Wahlgren et al., 2022). Additionally, longitudinal assessment across recovery is needed to assess the consequences of COVID-19 over time (Vanderlind et al., 2021).

Our aim was to assess subjective and objective cognitive dysfunction up to six months after COVID-19 in three different levels of acute disease severity: patients treated in the ICU or regular hospital wards or isolated at home. We examined the prevalence of subjective and objective cognitive impairment and the associations of subjective cognitive impairment to objective cognition, symptoms of depression and PTSD, and demographic and disease-related factors at three and six months after COVID-19.

## 2. Materials and methods

### 2.1. Study design, setting, and population

This is a post hoc study of the RECOVID project, a multidisciplinary,

single-center, prospective cohort study. The study protocol has been previously described (Ollila et al., 2022). The study is registered at [ClinicalTrials.gov](https://clinicaltrials.gov) (NCT04864938). The Ethics Committee of Helsinki University Hospital (HUS-1949-2020) approved the protocol. All participants gave written informed consent to the study and the principles of the Declaration of Helsinki were followed.

We assessed COVID-19 patients in three groups of acute disease severity (Fig. 1). A control group who reported no history of COVID-19 was also recruited (CONTROL group). The assessments were conducted three and six months after hospital discharge (ICU and WARD groups) or positive PCR test (HOME group). We included patients with data from at least one cognitive measure in at least one time point, resulting in a final study sample of 184 patients: 82 treated in ICU, 53 hospitalized who received ward-level care (WARD), and 49 home-isolated patients (HOME). The patients had been hospitalized or tested positive for COVID-19 between March 13 and December 31, 2020. Patients were typically admitted to the ICU if they had respiratory rate 30 and oxygen saturation with pulse oximeter 92% or below, were rapidly deteriorating with obviously increased work of breathing, or had other organ dysfunction necessitating intensive support or monitoring. The ICU admission criteria were in broad accordance with the published criteria for severe critical COVID-19 (Alhazzani et al., 2021). Recruitment took place by mailed invitation after discharge (ICU), from a pulmonary outpatient clinic appointment or during acute care in a pulmonology ward (WARD), or by media announcements (HOME and CONTROL). Inclusion criteria included a positive acute phase PCR or antibody test, age 18 or older, and Finnish as primary language. Exclusion criteria were pregnancy, prior major neurological diagnosis, developmental intellectual disability, or severely impaired hearing or vision.

### 2.2. Study outcomes

The main outcomes were the prevalence of subjective and objective cognitive impairment at 3 and 6 months after COVID-19 in three groups of acute disease severity compared to non-COVID controls. Secondary outcomes were the associations of subjective cognitive symptoms with objective cognitive impairment, demographic and disease-related factors, and depressive and PTSD symptoms.

We evaluated subjective cognitive impairment with the A-B Neuropsychological Assessment Schedule (ABNAS) questionnaire (Aldenkamp

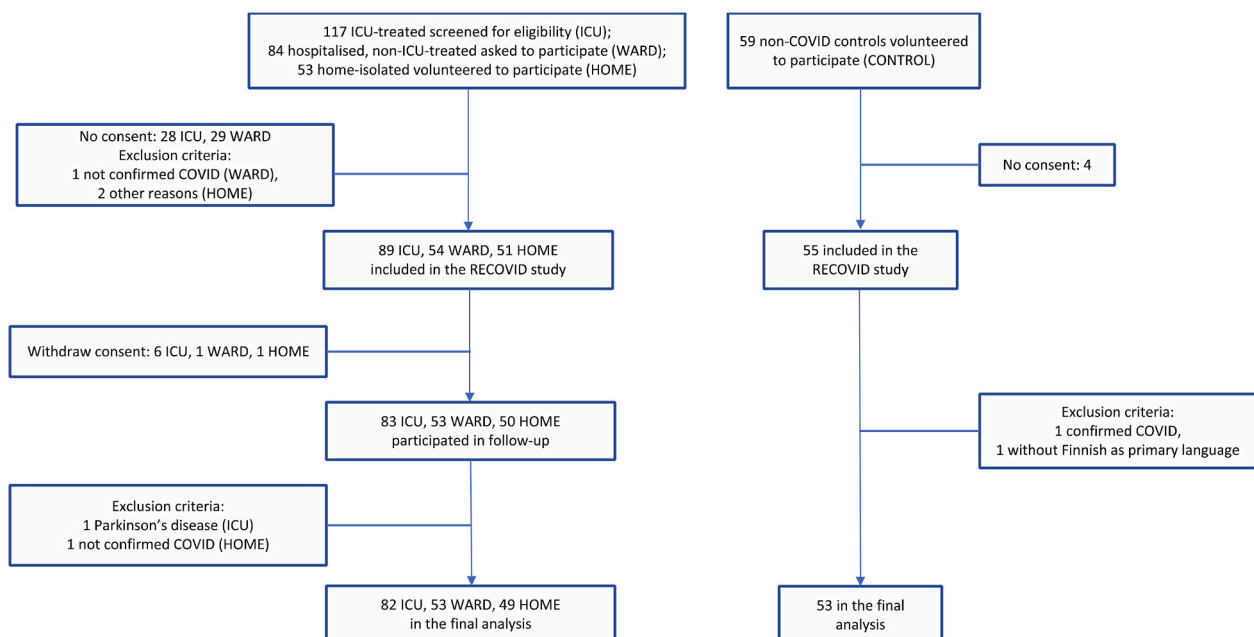


Fig. 1. Flow chart of the data compilation.

et al., 1995), a scale of patient-perceived cognitive functioning. It includes 24 items regarding cognitive impairments in daily life. Each item is scored on a scale from 0 to 3 with 0 indicating no difficulty, 1 mild difficulty, 2 moderate difficulty, and 3 severe difficulty. Cognitive symptoms are evaluated on six domains: fatigue (5 questions), slowing (5), memory (4), concentration (4), motor coordination (3), and language (3). The total score ranges from 0 to 72 with higher scores reflecting greater levels of subjective impairment. Aldenkamp et al. (2002) have proposed a dichotomization into low ( $\leq 15$ ) and high ( $> 15$ ) scores reflecting low and high levels of subjective cognitive impairment.

We screened objective cognitive functioning with Montreal Cognitive Assessment (MoCA). The telephone-based version (t-MoCA) (Katz et al., 2021) was used at 3 months. It includes a subset of items from the full MoCA, excluding items requiring visual stimuli and pencil-and-paper drawing. It consists of 22 items divided into domains of memory (5 points), attention (6 points), language (3 points), verbal reasoning (2 points), and orientation (6 points). In the 6-month assessment, we used the full version of MoCA, which has additional visuospatial/executive (5 points) and naming (3 points) domains. Total scores consist of points earned plus an additional point given with  $\leq 12$  years of education (no extra point is given with a perfect score) resulting in maximal total scores of 22 for t-MoCA and 30 for MoCA. Impaired MoCA scores were defined as  $\leq 19$  for t-MoCA (Pendlebury et al., 2013) and  $\leq 25$  for MoCA (Nasreddine et al., 2005). Experienced neuropsychologists or clinician investigators trained by experienced neuropsychologists performed the assessments.

We assessed post-traumatic stress disorder (PTSD) symptoms with the Impact of Event Scale-6 (IES-6), an abbreviated 6-item version of the IES-Revised (IES-R) that uses a five-point Likert scale ranging from 0 to 4 (Hosey et al., 2019). The score is formed as the mean of six items and the cut-off point indicating significant clinical symptoms of PTSD has been defined as 1.75 (Hosey et al., 2019).

We evaluated depressive symptoms with the two-question screening method (Depression screen) (Arroll et al., 2003; Whooley et al., 1997) at three months and the Patient Health Questionnaire-9 (PHQ-9) (Kroenke et al., 2001) at six months. The Depression screen was considered positive when “yes” was given to either of the two questions (Arroll et al., 2003). For the PHQ-9, a score of 10 or higher reflects at least moderate depressive symptoms (Kroenke et al., 2001).

We retrieved demographic information and clinical variables (length of hospital and ICU stay, supplementary oxygen, delirium, invasive mechanical ventilation [IMV], Charlson comorbidity index [CCI], and comorbidities) from medical records. Education level was reported by the patient.

### 2.3. Statistical analysis

We provide descriptive statistics for the demographic and clinical characteristics of each group and report the total sample size available for each data element. Sample size was based on power calculations for finding group differences which indicated that  $n > 50$  would be required for each group. Continuous variables are presented as mean (SD) or median (interquartile range [IQR]) and categorical variables as  $n$  (percentage). Group differences were assessed with ANOVA for normally distributed continuous variables, Kruskal-Wallis for non-normally distributed variables, and  $\chi^2$  test for categorical variables. Changes in the frequency of impairment in follow-up were assessed with McNemar tests. Because of the differences in age and education levels in the groups, we used additional age- and education-adjusted ANCOVAs with Tukey's post hoc tests to assess differences in MoCA scores. We transformed all ABNAS subscale scores to a range of 0–3 to enable comparison of subscales with differing numbers of items and assessed group differences with ANOVA.

We assessed the correlations of subjective cognitive symptoms with depressive and PTSD symptoms, demographic factors, hospital metrics, and objective cognitive impairment with Pearson's or Spearman's  $r$

correlations. We then used partial Pearson correlation coefficients controlling for age, education, and IES-6 to assess adjusted associations between subjective cognitive symptoms and objective cognitive performance. Finally, we used Venn diagrams to illustrate the overlap of objective cognitive dysfunction, subjective cognitive symptoms, and PTSD symptoms.

Statistical significance was set at  $p < .05$  and all tests were two-tailed. The Bonferroni correction and the Benjamini-Hochberg adjusted false discovery rate (FDR) were used to control for multiple comparisons. No imputation was performed for missing data. Data were analyzed using IBM SPSS Statistics for Mac Version 27 (IBM Corp., Armonk, NY) and Jamovi version 2.2 (The jamovi project, 2021, <https://jamovi.org>).

## 3. Results

### 3.1. Patient characteristics

The mean age of the COVID-19 patients was 53.4 (SD 12.8) years and 44% were men. The HOME group was younger than the ICU, WARD, and CONTROL groups (all  $p < .001$ ). The ICU group had less education than the WARD ( $p = .002$ ), HOME ( $p < .001$ ), and CONTROL ( $p < .001$ ) groups. The CONTROL group had 50.9% men and a mean age of 54.9 (12.3) years. Demographic characteristics, comorbidities, and hospital metrics are presented in Table 1.

### 3.2. Subjective cognitive symptoms

Of all COVID-19 patients, 40.6% (63/155) reported high levels of subjective cognitive symptoms (ABNAS score  $> 15$ ) at three months and 34.2% (53/155) at six months compared to 8.0% (4/50) of the CONTROL group. The frequencies of high ABNAS scores remained the same at three and six months (Table 2). There were no differences in ABNAS median scores between the ICU, WARD, and HOME groups (Table 2). In ABNAS subscales of fatigue ( $p < .001$ ) and language ( $p < .001$ ), all patient groups reported more symptoms than controls (Fig. 2). In slowing, more symptoms were reported by the ICU and HOME groups at 3 and 6 months and the WARD group at 6 months compared to controls. The ICU group reported more symptoms than controls in motor coordination at 3 and 6 months and the HOME group in concentration at 6 months.

### 3.3. Objective cognitive outcome

The 3-month cognitive assessment took place on average 108 (SD 18) days after hospital discharge for ICU and 152 (41) days for WARD patients and 181 (29) days after positive COVID-19 test result for HOME patients. Cognitive impairment was observed in 77 of 151 (51.0%) patients (Table 2). A statistically significant difference in the mean t-MoCA scores emerged between ICU (19.1 [95% CI 18.5 to 19.6]), WARD (18.7 [18.0 to 19.3]), and HOME (20.2 [19.7 to 20.7]) groups ( $p = .011$ ,  $\eta^2 0.06$ ). The HOME group scores were higher than ICU ( $p = .042$ ) and WARD ( $p = .009$ ) groups. Age ( $p = .015$ ,  $\eta^2 0.04$ ) and education ( $p < .001$ ,  $\eta^2 0.143$ ) were significantly related to t-MoCA score. After adjusting for age and education, a significant group effect remained ( $p = .038$ ,  $\eta^2 0.044$ ) but no significant post hoc pairwise differences were found after adjusting for multiple comparisons.

The 6-month cognitive assessment occurred on average 182 (18) days after hospital discharge for ICU and 198 (17) days for WARD patients and 220 (34) days after positive COVID-19 test result for HOME patients. Cognitive impairment was observed in 47 of 166 (28.3%) patients and 15 of 48 (31.3%) controls (Table 2). The frequency of objective cognitive impairment decreased in all COVID-19 groups from 3 to 6 months (Table 2). At 6 months, there was a statistically significant difference in the mean MoCA scores between ICU (26.4 [25.7 to 27.0]), WARD (26.3 [25.7 to 26.8]), HOME (28.2 [27.6 to 28.7]), and CONTROL (26.5 [25.8 to 27.2]) groups ( $p < .001$ ,  $\eta^2 0.096$ ). In the HOME

**Table 1**  
Demographic characteristics, comorbidities, and hospital metrics in COVID-19 patients and controls.

	ICU (n = 82)	WARD (n = 53)	HOME (n = 49)	$p^a$	All COVID-19 (N = 184)	CONTROL (n = 53)	$p^b$
<b>Demographic characteristics</b>							
Age, mean (SD), y	57.5 (11.6)	55.6 (9.6)	44.2 (13.3)	<.001	53.4 (12.8)	54.9 (12.3)	.453
Sex, male, No./total (%)	49/82 (59.8)	19/53 (35.8)	13/49 (26.5)	<.001	81/184 (44.0)	27/53 (50.9)	.373
Education, median (IQR), y	14 (12–15)	15 (13–17)	15 (14–17)	<.001	15 (12–17)	15 (13–17)	.115
<b>Comorbidities No./total (%)<sup>c</sup></b>							
CCI, median (IQR)	2 (1–3) n = 81	1 (1–2)	–	.196	–	–	–
Hypertension	45/82 (54.9)	15/53 (28.3)	8/48 (16.7)	<.001	68/183 (37.2)	11/53 (20.8)	.026
Hypercholesterolemia	26/82 (31.7)	11/53 (20.8)	4/48 (8.3)	.008	41/183 (22.4)	7/53 (13.2)	.143
Heart disease	12/82 (14.6)	3/53 (5.7)	5/48 (10.4)	.262	20/183 (10.9)	1/53 (1.9)	.042
Diabetes	19/82 (23.2)	5/53 (9.4)	2/48 (4.2)	.006	26/183 (14.2)	1/53 (1.9)	.013
Malignancy	6/82 (7.3)	3/53 (5.7)	2/48 (4.2)	.760	11/183 (6.0)	1/53 (1.9)	.229
Asthma	14/82 (17.1)	14/53 (26.4)	5/48 (10.4)	.108	33/183 (18.0)	3/53 (5.7)	.027
COPD	0/82 (0.0)	3/53 (5.7)	0/48 (0.0)	.024	3/183 (1.6)	0/53 (0.0)	.348
Kidney disease	3/82 (3.7)	0/53 (0.0)	0/48 (0.0)	.153	3/183 (1.6)	0/53 (0.0)	.348
Liver disease	0/82 (0.0)	2/53 (3.8)	2/48 (4.2)	.188	4/183 (2.2)	0/53 (0.0)	.278
<b>Hospital metrics</b>							
Delirium, No./total (%)	29/82 (35.4)	–	–	–	–	–	–
Duration of hospital stay, median (IQR), d	20 (15–26)	8 (5–11)	–	<.001	–	–	–
Duration of supplementary oxygen, median (IQR), d	20 (14–24) n = 78	5 (1–9)	–	<.001	–	–	–
Duration of ICU stay, median (IQR), d	10.5 (6–18)	–	–	–	–	–	–
Received IMV, No./total (%)	53/82 (64.6)	–	–	–	–	–	–
Duration of IMV, median (IQR), d <sup>d</sup>	7 (0–14)	–	–	–	–	–	–

Abbreviations: CCI, Charlson Comorbidity Index; COPD, Chronic Obstructive Pulmonary Disease; IMV, Invasive mechanical ventilation.

<sup>a</sup> ICU vs WARD vs HOME using ANOVA, Kruskal-Wallis test, or  $\chi^2$  test, as appropriate.

<sup>b</sup> Complete COVID-19 group vs non-COVID controls using *t*-test, Mann-Whitney test, or  $\chi^2$  test, as appropriate.

<sup>c</sup> Data missing from one patient in the HOME group.

<sup>d</sup> The duration of IMV in days included intubation and extubation days, and all episodes in the case of several IMV episodes.

group, scores were higher than in ICU ( $p < .001$ ), WARD ( $p < .001$ ), and CONTROL ( $p = .003$ ) groups. Age ( $p = .039$ ,  $\eta^2 0.02$ ) and education ( $p < .001$ ,  $\eta^2 0.171$ ) were significantly related to MoCA score. After adjusting for age and education, differences in MoCA scores were statistically significant ( $p = .006$ ,  $\eta^2 0.058$ ) with the HOME group having higher scores than the WARD ( $p = .006$ ) and CONTROL ( $p = .024$ ) groups.

### 3.4. Associations of subjective cognitive symptoms with psychological symptoms, demographic factors, hospital metrics, and objective cognitive impairment

IES-6 scores of  $>1.75$  were found in 27.7% of patients at 3 months and 19.6% at 6 months (Table 2). IES-6 scores correlated with ABNAS in all COVID-19 groups at six months and in the ICU and WARD groups at three months (Table 3). On the Depression screen at three months, 36.8% of COVID-19 patients were classified as being at risk of depressive symptoms (Table 2). PHQ-9 scores of  $\geq 10$  were discovered in 15.5% of COVID-19 patients at 6 months and 2.0% of controls (Table 2). Both depression scores correlated with ABNAS scores in all COVID-19 groups (Table 3).

Age and education did not correlate with ABNAS scores (Table 3). Sex was associated with ABNAS in COVID-19 patients: women (median 14 [IQR 6–27]) reported more subjective cognitive symptoms than men (median 10 [4–17.75]) at three months ( $p = .034$ ). A statistically significant difference was detected in the ABNAS score in the ICU group (women: median 15 [7.75–34.5]; men: median 10 [4.25–17];  $p = .018$ ) but not in the WARD or HOME groups. At six months, there were no sex differences in ABNAS scores in patient groups or controls.

No correlations were found between the length of hospital stay and ABNAS. In ICU patients, delirium (patients with and without delirium median 10.50 [5.75–28.5] and 11.50 [5.25–19.5], respectively,  $p = .431$ ) and IMV (with and without IMV median 12 [5–25] and 11 [7–23], respectively,  $p = .784$ ) were not associated with ABNAS.

There was a correlation with  $p = .05$  between MoCA and ABNAS at 3 months in the ICU group (Table 3). Because of strong associations between ABNAS and IES-6, additional partial correlation analyses

controlling for age, education, and IES-6 were performed. In the ICU group, the correlation between ABNAS and MoCA 3-month scores was not statistically significant after controlling for age, education, and IES-6 3-month score ( $r = -0.222$ ,  $p = .065$ ). No statistically significant partial correlations between MoCA and ABNAS were found at 3 or 6 months in the ICU, WARD, or HOME groups, or in the group of all COVID-19 patients combined.

The proportions of patients with significant subjective and objective cognitive symptoms and PTSD symptoms are described in Venn diagrams (Fig. 3).

## 4. Discussion

In this prospective single-center study of 184 COVID-19 patients and 53 controls we observed a poor congruence of subjective and objective cognitive function test results and an association of subjective cognitive impairment with depressive and PTSD symptoms in a six-month follow-up. Over one-third of patients with COVID-19, regardless of the acute disease severity, reported high levels of subjective cognitive impairment in a six-month follow-up. Objective cognitive impairment was observed in more than half of ICU and WARD patients and one-third of HOME patients at 3 months. At 6 months the cognitive performance of all three COVID-19 groups was similar (ICU and WARD) or better (HOME) than non-COVID controls. The frequency of objective but not subjective cognitive impairment decreased from 3 to 6 months. Depressive and PTSD symptoms were equally reported by all COVID-19 groups. Subjective cognitive symptoms were associated with depressive and PTSD symptoms, and in the ICU group, female sex. No associations were found between subjective cognitive impairment and objective cognitive impairment, length of hospital stay, delirium, IMV, age, or education.

The proportion of high levels of subjective cognitive symptoms in our study was approximately 30–40%, similar across the severity of acute disease, and stable in follow-up. Our findings differ from some studies where patients with more severe COVID-19 have reported more subjective symptoms (Pilotto et al., 2021; Halpin et al., 2021) but are in line with others which discovered similar rates of subjective cognitive



**Table 2**  
Cognitive and psychological outcomes in COVID-19 patients and controls.

	ICU (n = 82)	WARD (n = 53)	HOME (n = 49)	p	All COVID-19 (N = 184)	CONTROL <sup>a</sup> (n = 53)	p
<b>Subjective cognitive outcome</b>							
ABNAS score, 3 months, median (IQR)	11 (6–23.5) n = 74	11.5 (2–27) n = 46	14 (4–23) n = 35	.845	11 (4–24) n = 155	–	–
ABNAS score, 6 months, median (IQR)	10.5 (5–17) n = 62	10 (3–19) n = 51	14 (5–22) n = 42	.616	11 (4–19) n = 155	4.5 (2–9) n = 50	<.001
ABNAS >15, 3 months, No./total (%)	28/74 (37.8)	19/46 (41.3)	16/35 (45.7)	.732	63/155 (40.6)	–	–
[95% CI]	[26.8–49.9]	[27.0–56.8]	[28.8–63.4]		[32.8–48.8]		
ABNAS >15, 6 months, No./total (%)	20/62 (32.3)	19/51 (37.3)	14/42 (33.3)	.848	53/155 (34.2)	4/50 (8.0)	<.001
[95% CI]	[20.9–45.3]	[24.1–51.9]	[19.6–49.5]		[26.8–42.2]	[2.2–19.2]	
Difference in frequency of ABNAS >15, 3 vs 6 mo, p	.607	.774	.070	–	–	–	–
<b>Objective cognitive outcome</b>							
t-MoCA total score, 3 months, median (IQR)	19 (17–21) n = 77	19 (17–20) n = 44	20 (19–21) n = 30	.011	19 (18–21) n = 151	–	–
MoCA total score, 6 months, median (IQR)	27 (25–29) n = 72	26 (25–28) n = 49	29 (27–30) n = 45	<.001	27 (25–29) n = 166	27 (25–28) n = 48	.366
t-MoCA ≤19, 3 months, No./total (%)	41/77 (53.2)	27/44 (61.4)	9/30 (30.0)	.025	77/151 (51.0)	–	–
[95% CI]	[41.5–64.7]	[45.5–75.6]	[14.7–49.4]		[42.7–59.2]		
MoCA ≤25, 6 months, No./total (%)	26/72 (36.1)	17/49 (34.7)	4/45 (8.9)	.003	47/166 (28.3)	15/48 (31.3)	.693
[95% CI]	[25.1–48.3]	[21.7–49.6]	[2.5–21.2]		[21.6–35.8]	[18.7–46.3]	
Difference in frequency of impaired MoCA scores, 3 vs 6 months, p	.007	.003	.008	–	–	–	–
<b>Depressive symptoms</b>							
Depression screen ≥1, 3 months, No./total (%) [95% CI]	29/74 (39.2) [28.0–51.2]	18/46 (39.1) [25.1–54.6]	10/35 (28.6) [14.6–46.3]	.520	57/155 (36.8) [29.2–44.9]	12/48 (25) [13.6–39.6]	.132
PHQ-9 ≥ 10, 6 months, No./total (%) [95% CI]	8/62 (12.9) [5.7–23.9]	8/51 (15.7) [7.0–28.6]	8/42 (19.0) [8.6–34.1]	.696	24/155 (15.5) [10.2–22.2]	1/50 (2.0) [0.1–10.6]	.011
<b>PTSD symptoms</b>							
IES-6 > 1.75, 3 months, No./total (%) [95% CI]	24/74 (32.4) [22.0–44.3]	13/46 (28.3) [16.0–43.5]	6/35 (17.1) [6.6–33.6]	.249	43/155 (27.7) [20.9–35.5]	–	–
IES-6 > 1.75, 6 months, No./total (%) [95% CI]	13/62 (21.0) [11.7–33.2]	10/49 (20.4) [10.2–34.3]	7/42 (16.7) [7.0–31.4]	.851	30/153 (19.6) [13.6–26.8]	2/49 (4.1) [0.5–14.0]	.010
Difference in frequency of IES-6 > 1.75, 3 vs 6 months, p	.109	.344	.687	–	–	–	–

Abbreviations: ABNAS, A-B Neuropsychological Assessment Schedule; MoCA, Montreal Cognitive Assessment; PHQ-9, Patient Health Questionnaire-9; IES-6, Impact of Event Scale-6.

<sup>b</sup> McNemar test p values.

<sup>a</sup> Data collected once for the CONTROL group.

deficits regardless of disease severity (Blackmon et al., 2022; Rass et al., 2022). Moreover, our results strengthen the finding that COVID-19 patients not requiring hospitalization can also manifest subjective cognitive symptoms (Vanderlind et al., 2021).

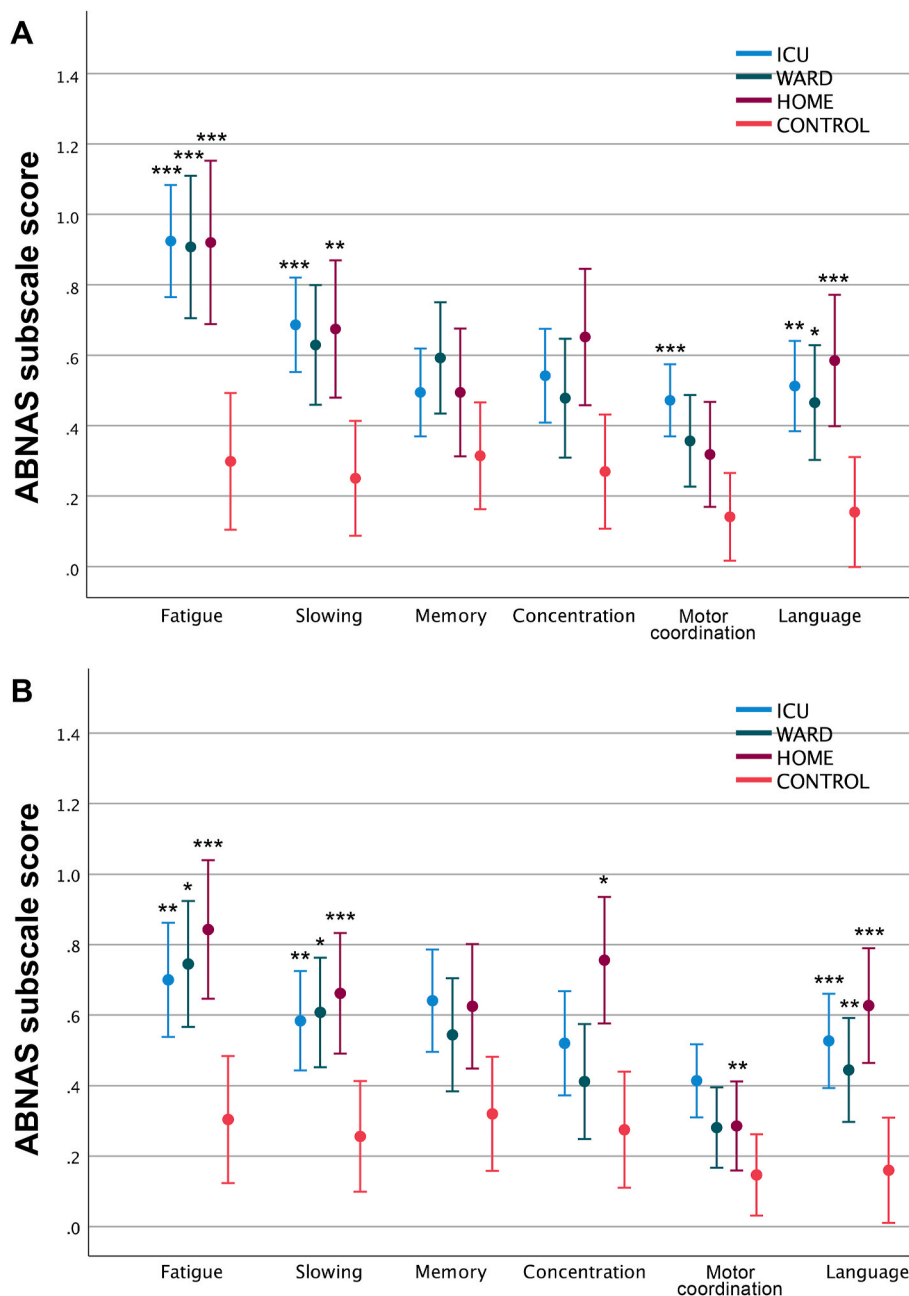
COVID-19 patients reported more subjective cognitive symptoms than controls especially in the domains of fatigue, slowing, and language. Fatigue has been one of the most reported symptoms after COVID-19 (Graham et al., 2021). An interesting finding was the high amount of subjective language symptoms among COVID-19 patients, which warrants further research. In the concentration domain, the HOME group reported the most symptoms, which may reflect larger demands in work and daily life as they were the youngest group. Problems in motor coordination, on the other hand, were reported by the ICU group. Muscular weakness and decreased mobility are known symptoms of post-intensive care syndrome (Needham et al., 2012). Interestingly, COVID-19 patients' subjective symptoms in memory did not differ from non-COVID controls. Blackmon et al. (2022) reported that 30–40% of COVID-19 patients showed subjective memory and attention problems, but they had no data on controls. This highlights the importance of a control group in COVID-19 studies. During a pandemic, even healthy individuals without coronavirus disease may experience subjective difficulties relating to the overall burden of the pandemic (e.g., restrictions and social isolation) or other, unexamined factors. Given the observed psychological distress in the general population during the pandemic (Patel et al., 2022), problems in cognitive functioning are also possible.

In objective cognitive screening at three months, more than half of ICU and WARD and almost one-third of HOME patients had t-MoCA

scores in the impaired range. The percentages of impaired MoCA scores decreased in follow-up. At six months, the rates of impaired cognitive performance in the ICU and WARD groups correspond to the findings of previous studies of hospitalized COVID-19 patients (Morin et al., 2021; Nersesjan et al., 2022; Raman et al., 2021). However, in our study, a similar proportion of the non-COVID controls also had MoCA scores in the impaired range. It is noteworthy that our non-COVID control sample consisted of individuals with no acute illness, whereas some studies have included patients hospitalized for other diseases as control subjects (Nersesjan et al., 2022).

Subjective cognitive symptoms were not associated with objective cognitive screening, length of hospital stay, delirium, or IMV. Some previous studies have found subjective cognitive complaints to correlate with objectively measured cognitive impairment (Miskowiak et al., 2021, 2022) whereas others (Almeria et al., 2020; Costas-Carrera et al., 2022) have not. Previous studies have reported higher incidences of either objective (Ceban et al., 2022; Mendez et al., 2022) or subjective cognitive dysfunction (Venturelli et al., 2021). Assessment methods and the definitions of impairment vary across studies. Self-reported symptoms do not necessarily manifest as impaired performance in neuropsychological tests, especially in a short cognitive screen as used in the present study. Unmeasured factors may also affect cognitive assessment (Valdes et al., 2022).

Psychological factors may also have a role in persisting subjective cognitive complaints after COVID-19 (Krishnan et al., 2022). In our study, subjective cognitive complaints were associated with depressive and PTSD symptoms, which was expected since psychological factors have been shown to associate with subjective cognitive complaints



**Fig. 2.** Mean ABNAS subscale scores at a) three and b) six months after COVID-19. The subscale scores were transformed to a range of 0–3 to enable comparison of subscales with differing numbers of items. Control group scores are shown in both time points. Error bars represent 95% confidence intervals. Bonferroni adjusted p values are reported. \* $p < .05$ , \*\* $p < .001$ , \*\*\* $p < .001$  compared to CONTROL. (Color: online only.). (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

(Almeria et al., 2020; Brown et al., 2022; Henneghan et al., 2022; Liyanage-Don et al., 2022). The relationship between disease-related characteristics and subjective cognitive symptoms is still unclear. Consistent with our results, disease-related characteristics were not associated with self-reported fatigue in a previous study (Townsend et al., 2020). However, a greater number of COVID-related symptoms during the infection correlated with more subjective cognitive problems in another study (Henneghan et al., 2022). In our study, women in the ICU group reported more subjective cognitive symptoms than men. Female sex has previously been reported to be associated with incomplete recovery (Evans et al., 2021), more frequent subjective cognitive decline (Ferrucci et al., 2021; Liyanage-Don et al., 2022), and more severe fatigue after COVID-19 (Halpin et al., 2021; Townsend et al., 2020).

The strengths of our study include using a non-COVID control group which allows comparison to individuals who have been exposed to the indirect, societal factors associated with the pandemic. This is one of the first studies, to our knowledge, to investigate objective and subjective

cognitive outcomes in several levels of COVID-19 severity. Additionally, the multidisciplinary approach allowed a comprehensive examination of clinical, cognitive, and psychological factors.

There are several limitations to this study. First, the home-isolated group was the youngest and predominantly female, whereas the ICU group was predominantly male with the lowest education level, reflecting the risk factors for severe COVID-19 infection. There is a rather large sex difference in the home-isolated group with a high number of female subjects. In previous research, females have reported more long-term COVID-19 symptoms than males, which may have several possible explanations such as sex-related immune system differences or gender-related social factors (Pela et al., 2022). We cannot exclude selection bias, inherent to clinical research. Second, the MoCA is a cognitive screening instrument with limited sensitivity to detect subtle cognitive deficits in younger, non-neurodegenerative populations such as our cohort (Ceban et al., 2022). In our earlier study, we found impairment especially in the ICU-treated group in domains of attention

**Table 3**  
Correlations of ABNAS scores to cognitive, psychological, demographic, and admission variables.

	ICU		WARD		HOME		CONTROL
	3 months <sup>a</sup>	6 months <sup>b</sup>	3 months <sup>c</sup>	6 months <sup>d</sup>	3 months <sup>e</sup>	6 months <sup>f</sup>	ABNAS <sup>g</sup>
ABNAS, 6 months	.792 *** n = 59	–	.669 ***	–	.820 *** n = 31	–	–
t-MoCA, 3 months	–.253 <sup>†</sup> n = 73	–.179 n = 58	–.151 n = 41	–.238 n = 43	–.014 n = 26	.034 n = 28	–
MoCA, 6 months	–.051 n = 65	.041 n = 58	–.100 n = 43	–.114 n = 47	.079 n = 32	–.106 n = 40	–.081 n = 45
Depression screen, 3 months	.519 ***	.338 * n = 59	.587 ***	.491 ** n = 46	.439 *	.263 n = 31	.411 ** n = 48
PHQ-9, 6 months	.590 *** n = 59	.816 ***	.651 ***	.838 ***	.653 *** n = 31	.825 ***	.686 ***
IES-6, 3 months	.642 ***	.540 *** n = 59	.599 ***	.388 * n = 46	.379 *	.256 n = 31	–
IES-6, 6 months	.517 *** n = 59	.653 ***	.379 * n = 45	.557 *** n = 49	.534 ** n = 31	.657 ***	.400 ** n = 49
Length of hospital stay	–.040	–.044	.017	–.192	–	–	–
Age	.097	–.015	–.053	–.032	–.003	–.248	–.216
Education	–.071	.189 n = 61	–.191 n = 45	–.190 n = 50	.162	–.213	–.158

\* $p < .05$ , \*\* $p < .001$ , \*\*\* $p < .001$ . <sup>†</sup> $p = .05$ . FDR-adjusted p-values are reported.

<sup>a</sup> n = 74.

<sup>b</sup> n = 62.

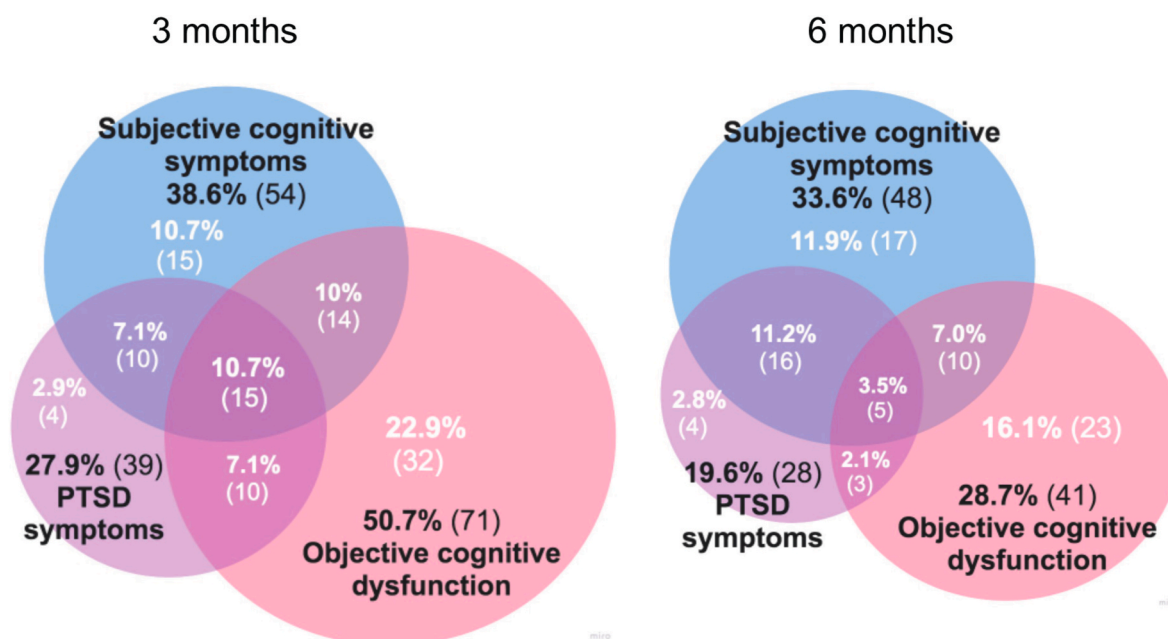
<sup>c</sup> n = 46.

<sup>d</sup> n = 51.

<sup>e</sup> n = 35.

<sup>f</sup> n = 42.

<sup>g</sup> n = 50 unless otherwise specified.



**Fig. 3.** Venn diagrams of the overlap of subjective cognitive symptoms, objective cognitive dysfunction, and PTSD symptoms. Subjective cognitive symptoms were defined as an ABNAS score >15, objective cognitive dysfunction as a t-MoCA score ≤19 (3 months) and a MoCA score ≤25 (6 months), and significant PTSD symptoms as an IES-6 mean score >1.75. The 3-month (n = 140) and 6-month (n = 143) follow-ups are shown. (Color: online only.). (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

and executive functions, which are difficult to evaluate with a screening test (Ollila et al., 2022). We used both the telephone and in-person versions of the MoCA, which limits the comparability of the results. Third, there were missing data during follow-up especially for the HOME group. Fourth, we cannot exclude volunteer bias, as subjects concerned about their cognition may have been more interested in participating the study. Fifth, we did not use an antibody test to confirm that the controls had not had an asymptomatic COVID-19 infection. Thus, we cannot fully exclude the possibility of a previous asymptomatic infection in the non-COVID controls. Sixth, to our knowledge, the ABNAS was used for the first time to assess subjective cognitive functioning after COVID-19. However, it is one of the methods recommended by the NEUROCovid Neuropsychology Taskforce (Cysique et al., 2021). Finally, this is a single-center study which limits the generalizability of

our results.

### 5. Conclusions

In conclusion, long-term subjective cognitive impairment and psychological symptoms were present in COVID-19 patients regardless of the severity of the acute disease. Subjective cognitive symptoms were associated with depressive and PTSD symptoms and female sex, but not with objective cognitive screening. Objective neuropsychological assessment and self-reported measures of cognition seem to assess different constructs, although providing valuable information on different aspects of functioning. It is important to thoroughly assess patients reporting long-term subjective symptoms and screen for underlying mental health related factors such as depression or PTSD.



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## Declaration of competing interest

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## Data availability

Data not publicly available, because consent for public sharing was not included in the original consent formulation. Data can be partially shared within the EU/EESC upon request for research purposes

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