Additional File 1. Sequelae, persistent symptomatology, and outcomes after COVID-19 hospitalization: the ANCOHVID multicentre 6-month follow-up study.

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## **STROBE Statement**

Checklist of items included in reports of observational studies.

	Item No	Recommendation
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title
		or the abstract. Retrospective observational longitudinal (6-month
		follow-up) study (abstract).
		(b) Provide in the abstract an informative and balanced summary of
		what was done and what was found. Provided in the abstract.
Introduction		
Background/rationale	2	Explain the scientific background and rationale for the investigation
		being reported. Explained in the first 4 paragraphs of the
		introduction.
Objectives	3	State specific objectives, including any prespecified hypotheses.
		Stated in the last (fifth) paragraph of the introduction.
Methods		
Study design	4	Present key elements of study design early in the paper. <b>Presented at</b>
, ,		the beginning of the methods (study design and setting).
Setting	5	Describe the setting, locations, and relevant dates, including periods
		of recruitment, exposure, follow-up, and data collection. Described
		at the beginning of the methods (study design and setting).
Participants	6	Give the eligibility criteria, and the sources and methods of selection
		of participants. Presented at the beginning of the methods (study
		design and setting).
Variables	7	Clearly define all outcomes, exposures, predictors, potential
		confounders, and effect modifiers. Give diagnostic criteria, if
		applicable. We presented all the variables of the study in the "data
		source and variables" section. We indicated the exposure and
		outcome variables.
Data sources/	8*	For each variable of interest, give sources of data and details of
measurement		methods of assessment (measurement). Describe comparability of
		assessment methods if there is more than one group. We presented
		all the sources of the variables of the study in the "data source
		and variables" section and provided a full description in

		Additional File 2: Table S1. We also performed a dendrogram for		
		cluster analysis (figure 1) in order to provide evidence of validity		
D'	0	of the information collected.		
Bias	9	Describe any efforts to address potential sources of bias. <b>Described</b>		
a. 1 .	10	in "statistical analyses" section.		
Study size	10	Explain how the study size was arrived at. <b>Explained in the methods</b>		
		(a complete cohort of hospitalized patients during the period of		
		recruitment).		
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If		
		applicable, describe which groupings were chosen and why.		
		Explained in the methods.		
Statistical methods	12	(a) Describe all statistical methods, including those used to control		
		for confounding. Described in "statistical analyses" section.		
		(b) Describe any methods used to examine subgroups and		
		interactions. Described in "statistical analyses" section.		
		(c) Explain how missing data were addressed. Explained throughout		
		the manuscript (two variables were not considered for presenting		
		too many missing values: smoking habit and obesity).		
		If applicable, describe analytical methods taking account of sampling		
		strategy. N/A.		
		(e) Describe any sensitivity analyses. N/A.		
Participants 13*	(a) Report numbers of individuals at each stage of study—eg numbers potentially eligible, examined for eligibility, confirmed eligible, included in the study, completing follow-up, and analysed. <b>Explained in the first</b>			
	paragraph of the "Sociodemographic and clinical variables" section.			
	(b) Gi	ive reasons for non-participation at each stage. We included all		
	hospi	talized patients during recruitment (non-participation is not		
	applic	cable in this study).		
	(c) Consider use of a flow diagram. N/A			
Descriptive 14*	(a) Give characteristics of study participants (eg demographic, clinical, social)			
data	and information on exposures and potential confounders. <b>Explained in the first paragraph of the "Sociodemographic and clinical variables" section and in Table 1.</b>			
	and n	n Table 1.		
		dicate number of participants with missing data for each variable of st. Indicated in Table 1 ("Unknown").		
	111010	on and a rest of Chimoth ).		
Outcome data 15*	Cross-sectional study—Report numbers of outcome events or summary measures. Reported in Table 1 (columns).			
Main results 16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted estimates and their precision (eg, 95% confidence interval). Make clear which confounders were adjusted for and why they were included. <b>Given in Additional File 2: Tables S3 to S5.</b>			

		(b) Report category boundaries when continuous variables were categorized. <b>Reported in Table 1 (age).</b>		
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period. <b>N/A</b>		
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses. Reported in the "Frequency of SPS during the 6-month follow-up" and "Outcomes" sections.		
Discussion				
Key results	18	Summarise key results with reference to study objectives. <b>Reported in the first paragraph of the Discussion.</b>		
Limitations	19	Discuss limitations of the study, taking into account sources of potential bias or imprecision. Discuss both direction and magnitude of any potential bias. <b>Thoroughly reported in the Limitations subsection.</b>		
Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence. <b>Reported throughout the Discussion and Conclusion sections.</b>		
Generalisability	21	Discuss the generalisability (external validity) of the study results. <b>Reported</b> in the Limitations subsection		
Other informati	on			
Funding	22	Give the source of funding and the role of the funders for the present study and, if applicable, for the original study on which the present article is based. <b>Reported in the "Funding" declaration.</b>		