

INFORMED CONSENT TO PARTICIPATE IN THE STUDY:

"Pulmonary telerehabilitation for patients affected by COVID-19, confined to their homes: controlled and randomized clinical trial"

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Additional information and resolution of questions related to the study: cleofas@us.es

- I have read the patient study information sheet, hosted at this web address:

<https://www.fisiosurid.com/researchcovid19>

- I meet the inclusion criteria to participate in it, which are:

- You must be between 18-75 years old
- You must have symptoms consistent with COVID-19 after having close contact with a COVID-19 infected patient, or you have tested positive for coronavirus (COVID-19) polymerase chain reaction (PCR), or in the antigen test, from 40 days before the start of the study, or up to 10 days after it.
- I have been able to ask questions about the study, by talking to the researchers or email. (cleofas@us.es)

- I have been able to receive enough information about the study.

- I understand that my authorization is voluntary.

- I understand that I can withdraw from the study:

1. When I want, of my own free will.
2. Without having to explain my decision.
3. Without this affecting my health care.

- I freely grant my consent to participate in the study.

- I give my consent to participate in this study, and for this I will have to fill out and send the informed consent form to the researchers, hosted at this web address:

<https://www.fisiosurid.com/researchcovid19>