INFORMED CONSENT TO PARTICIPATE IN THE STUDY:

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

Head Researchers: Dr. Cleofás Rodríguez-Blanco - Dr. Juan José González Gerez
Research Group SICA, CTS954: Innovations in Health and Quality of Life.
Additional information and resolution of questions related to the study: cleofas@us.es

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

  https://www.fisiosurid.com/covid-19/

- 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.He podido hacer preguntas sobre el estudio, hablando con alguno/a de los investigadores/as, o bien a través del correo electrónico (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 you want, of your 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 principal investigator, hosted at this web address:

  https://www.fisiosurid.com/covid-19/