Informed consent to participate in a health science research study.

Title of the research project: “The role of hyperglycemia, hyperinsulinemia and elevated free fatty acids for cardiac function in patients with type 2 diabetes – the HyperCarD2 study”.

Declaration from the patient:

I have received written and oral information and I know enough about the purpose, method, benefits and disadvantages of saying yes to participating.

I know that participating is voluntary and that I can always withdraw my consent without losing my current or future rights to treatment.

I give my consent to participate in the research study and to have my biological material collected and stored in a research biobank. I have received a copy of this consent form as well as a copy of the written information about the study for my own use.

Name of the patient: ______________________________________________________

Date: _______________   Signature: ____________________________________________

If new essential health information about you appears in the research study, you will be informed. If you would like to decline receiving any new information about your health that appears in the research project, please mark here: __________ (insert an x)

Do you want to be informed about the result of the research study and any possible consequences for you?
Yes _____ (insert an x)         No _____ (insert an x)

Declaration from the person providing the information:

I declare that the patient has received oral and written information about the research study.

In my opinion, sufficient information has been provided to enable a decision to be taken on participation in the study.

The name of the person who provided the information: Roopameera Thirumathyam

Date: _______________   Signature: ____________________________________________

Study identification: (E.g. comité ID, EudraCT no., version no./date or similar.)

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