THE DANISH BIOMEDICAL RESEARCH ETHICS COMMITTEE SYSTEM

Standard consent statement prepared by the Danish Biomedical Research Ethics Committee System, December 2011. (S4)

Informed consent to participate in a health science research project

Title of the research project:

Can-Art

Pain treatment with medical CANnabis in patients with inflammatory ARThritis

Statement by Research Subject:

I have been given written and oral information and I know enough about the purpose and method, and about the advantages and disadvantages of participation.

I know that participation is voluntary and that I can withdraw my consent at any time without losing my current or future rights to treatment.

I consent to participate in the research project and that my biological data be extracted for storage in a research bio-bank. I have received a copy of this consent form and a copy of the written information about the project for my own records.

Research subject's name: _____

Date _____ Signature: _____

If new, essential health information about you comes to light in the research project, you will be informed. If you would prefer *not* to be informed about new, essential health information, should it come to light in the research project, please mark here: _ (mark with an x)

Do you want to be informed about the results of the research project and any consequences for you?

Yes _____ (mark with an x) No _____ (mark with an x)

Declaration by the person providing this information:

I declare that the research subject has received oral and written information about the research project.

In my opinion, sufficient information has been provided to the research subject for the decision to be made regarding participation in the research project.

The name of the person who provided this information:

Date _____Signature: _____

Project identification: (VEK Project ID 61 187, EUdraCT no. 2017-2017-004226-15)