Human Subject Research Projection

Name

Institution

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After watching the instructional videos, I, being research coordinator—Jan Klein—will not abide by my PI when he asks me to falsify study data and violate study protocol for the melanoma subjects. I will instantly talk to my supervisor about this issue and her timely intervention will definitely help me get rid of this ethical dilemma. Moreover, I will arrange weekly educational sessions to keep my research staff and PI updated about the newly emerging protocols within the research field. In this way, careful decision-making will help ensuring patient’s protection in research scenarios.

In order to protect the human subjects, informed consent is a mandatory paper to be signed having an explanation about the purpose of research, probable time-period of subject’s involvement and a fine delineation of procedures (tests, experimentation) to be followed. Informed consent will provide a brief fact-sheet to the subjects about study protocols and ensuring their privacy rights prior to undertaking research (Lokesh et. al., 2016).

Being a registered nurse (RN) and clinical research coordinator (CRC), it is quite essential for me to protect the human and legal rights of the patients and provide assistance in pronouncing those rights if need arises. I will make sure that the client is agreed upon receiving information and then I will present it in the best understandable way. Besides informal support, the objective support will include the right to choose values that seem necessary to maintain patient’s lives and the right to decide which scheme of actions will achieve the desired values (Vaartio et. al., 2006). In this way, a good decision making and profound insight about the updated research protocols will help me ensuring patients’ protection in the specific melanoma case.

References

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