

APPROVAL OF RESEARCH Using Expedited Procedures

March 08, 2018

Type of Review:	Submission Response for Initial Review Submission
Type of neview.	Form
Title:	Tracking Kinematic and Kinetic Data during Horse
Title:	Riding for Optimizing Therapeutic Outcomes
Investigator:	Pilwon Hur, PhD
IRB ID:	IRB2018-0064
Reference Number:	070687
Funding:	Horses and Humans Research Foundation(HHRF)
Documents Approved:	IRB Application 1.1
Bocaments Approved.	THE Application 1.1
	Child Assent 1.0
	Parent Permission Consent 1.0
	Human Research Subject Recruitment Script 1.0
Special Determinations:	(45 CFR 46.404/ 21 CFR 50.51): Not greater than
	minimal risk
	Assent waived under 45 CFR 46.408/ 46.116
	One parent signature
Risk Level of Study:	Not Greater than Minimal Risk under 45 CFR 46 / 21
,	CFR 56
Review Category:	Category 4: Collection of data through noninvasive
,	procedures (not involving general anesthesia or
	sedation) routinely employed in clinical practice,
	excluding procedures involving x-rays or microwaves.
	Where medical devices are employed, they must be
	cleared/approved for marketing
	Category 6: Collection of data from voice, video,
	digital, or image recordings made for research
	purposes

Dear Pilwon Hur, PhD:



The IRB approved this research from 03/08/2018 to 03/07/2019 inclusive.

It is recommended that you submit your next continuing review by 02/07/2019 to avoid a lapse in approval. Your study approval will end on 03/07/2019.

Your study must maintain an **approved status** as long as you are interacting or intervening with living individuals or their identifiable private information or identifiable specimens.

Obtaining identifiable private information or identifiable specimens includes, but is not limited to:

- 1. using, studying, or analyzing for research purposes identifiable private information or identifiable specimens that have been provided to investigators from any source; and
- 2. using, studying, or analyzing for research purposes identifiable private information or identifiable specimens that were already in the possession of the investigator.

In general, OHRP considers private information or specimens to be individually identifiable as defined at 45 CFR 46.102(f) when they can be linked to specific individuals by the investigator(s) either directly or indirectly through coding systems.

If you have any questions, please contact the IRB Administrative Office at 1-979-458-4067, toll free at 1-855-795-8636.

Sincerely, IRB Administration