

University of Illinois at Urbana–Champaign

Institutional Review Board Office

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WAIVER OF DOCUMENTATION OF INFORMED CONSENT (45CFR46.117(C))

ALL APPLICATIONS MUST BE TYPEWRITTEN, SIGNED, AND SUBMITTED AS SINGLE-SIDED HARD COPY. PLEASE, NO STAPLES!

Responsible Project Investigator (R	.PI):		
Last Name: Twidale		First Name: Michael B.	Dept. or Unit: Grad. Sch. of Lib.&Info. Sci
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Project Title:			
Investigating the data quality effect	of task	design changes in paid cro	wd systems
			sent, please provide a response to EITHER ner statement is true for this research.
would be potential harm resulting f wants documentation linking the su	rom a bi	reach of confidentiality. Ea ith the research, and the sul	the consent document and the principal risk ch subject will be asked whether the subject oject's wishes will govern. *Note: A waiver of egory if the research is subject to FDA
(2) The research presents no more to consent is normally required outside			ets and involves no procedures for which written
process will be conducted online and The study involves completing task would not be required outside of the	nd by co s that ar e researc	mpleting the tasks, participe not risky, so the study is ch context. For example, if	of informed consent from the IRB. The consent ants are implying their consent to participate. no more than minimal risks. Informed consent Amazon were collecting this data for quality orkers, informed consent would not be
** In cases in which the documenta subjects with a written statement re			B may require the investigator to provide
RPI Signature: MP Tuiled	2		Date: <u>2/4/15</u>
IRB Member Approval:			Date: