



## Office of Research Administration

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DATE: September 13, 2012

TO: Kristian Marlow, B.A.  
FROM: University of Missouri-St. Louis IRB

PROJECT TITLE: [373693-2] Application for Expedited Review of the Use of Human Subjects for a Deaf Motion-Sound Synesthete Study

REFERENCE #:  
SUBMISSION TYPE: Amendment/Modification

ACTION: APPROVED

APPROVAL DATE: September 13, 2012

EXPIRATION DATE: September 13, 2013

REVIEW TYPE: Expedited Review

REVIEW CATEGORY: Expedited review category # 7

The chairperson of the University of Missouri-St. Louis IRB has reviewed the above mentioned protocol for research involving human subjects and determined that the project qualifies for expedited review under Title 45 Code of Federal Regulations Part 46.110b. The time period for this approval expires one year from the date listed below. You must notify the University of Missouri-St. Louis IRB in advance of any proposed major changes in your approved protocol, e.g., addition of research sites or research instruments.

You must file an annual report with the committee. This report must indicate the starting date of the project and the number of subjects to date from start of project, or since last annual report, whichever is more recent.

Any consent or assent forms must be signed in duplicate and a copy provided to the subject. The principal investigator must retain the other copy of the signed consent form for at least three years following the completion of the research activity and they must be available for inspection if there is an official review of the UM-St. Louis human subjects research proceedings by the U.S. Department of Health and Human Services Office for Protection from Research Risks.

This action is officially recorded in the minutes of the committee.

If you have any questions, please contact Carl Bassi at 314-516-6029 or [bassi@umsl.edu](mailto:bassi@umsl.edu). Please include your project title and reference number in all correspondence with this committee.