**Office of Research Administration**

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| DATE: | November 14, 2012 |
| TO: | Matt DeStefano |
| FROM: | University of Missouri-St. Louis IRB |
| PROJECT TITLE: | [376281-2] Contrast and Grapheme-Color Synesthesia |
| REFERENCE #: | |
| SUBMISSION TYPE: | Revision |
| ACTION: | MODIFICATIONS APPROVED |
| DECISION DATE: | November 14, 2012 |
| EXPIRATION DATE: | November 14, 2013 |
| REVIEW TYPE: | Expedited Review |

This modification was approved by the University of Missouri-St. Louis IRB for the term of this protocol. The University of Missouri-St. Louis IRB must be notified in writing prior to major changes in the approved protocol. Examples of major changes are the addition of research sites or research instruments.

An annual report must be filed with the committee. This report should indicate the starting date of the project and the number of subjects since the start of project, or since last annual report.

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. Please include your project title and reference number in all correspondence with this committee.