Solutions Institutional Review Board
4925 Copper Creek Lane
Little Rock, AR 72223

Dear Amy Johnson,

You are hereby notified that the IRB APPROVED, by expedited review, the above referenced new study protocol on 02/20/2015. Your study may now commence.

Furthermore, your protocol is approved for a period of one year effective 02/20/2015 and ending 2/19/2016. This action will be reported to the full IRB on 03/02/2015.

Please be advised that it is required that a Continuing Review Form, be submitted two (2) months prior to the 2/19/2016 IRB meeting. If the study is completed before that date, a final Continuing Review Form, should be submitted at the close of the study.

Your study must be conducted in accordance with the Solutions Institutional Review Board IRB Policy and Procedures Manual and applicable law. Your responsibilities include but are not limited to:

- Changes to the protocol or its related informed consent document must be approved by the IRB prior to implementation.
- Adverse events must be reported promptly to the IRB.
- Each subject should receive a copy of the informed consent document.
- Records must be retained for at least three years.

Sincerely,
Dana Gonzales
Solutions Institutional Review Board