# **IRB Protocol Closure Form**

Human Participants in Non-medical Research

1. Were any anticipated problems encountered throughout the duration of the study that have not been reported to the IRB? If so, please describe.
2. Were any adverse events encountered throughout the duration of the study that have not been reported to the IRB? If so, please describe.
3. Did you deviate from the approved protocol or approved amendments? If so, please describe.