

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.