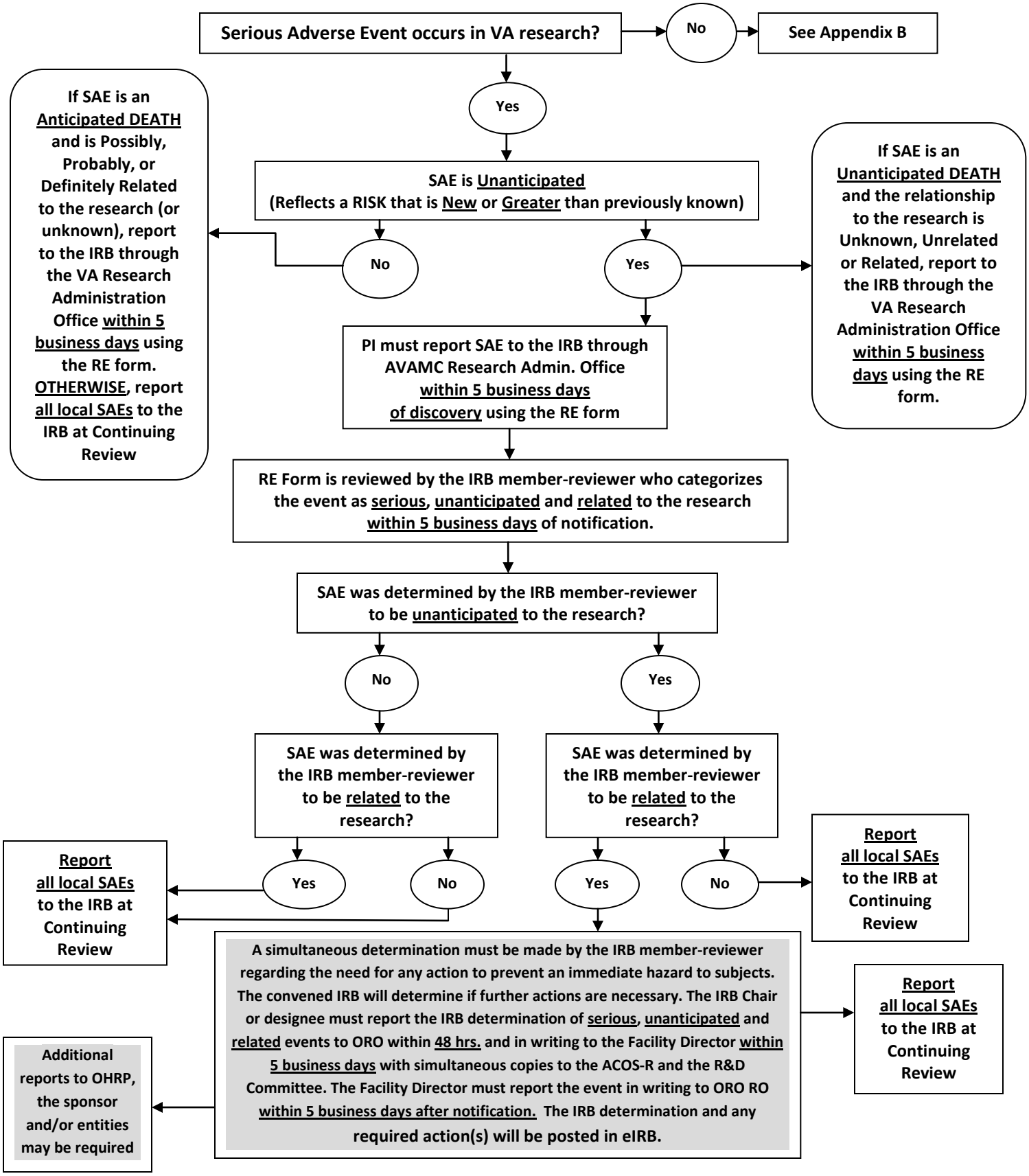
Appendix A
Reportable Events – Serious Adverse Events



*Shaded areas indicate reporting responsibility of the IRB/facility and not the responsibility of the study team.