

Appendix 5 – PHARMACIST

1. How are investigational drugs and devices stored? Temperature control? Access control?
2. Is there any differentiation between investigational drugs and regular clinical care drugs?
3. How do inpatients and outpatients receive investigational drugs and devices?
4. Is the pharmacy involved in preparation of investigational materials? Reconstitution? Preparation of placebos?
5. Do you have a mechanism to ensure that all clinical trials requesting for the pharmacy services have IRB approval?
6. How do you ensure that only clinical trial participants receive investigational materials?
7. After the end of the study, how are investigational materials disposed?
8. Do you have any message you would like us to convey to the University officials?