附表1

参加试验人员表

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| 研究职责 | | | | | | | | | | | |
| 1 | 获取知情同意书 | | 2 | | 受试者筛选 | | 3 | 受试者随访评估 | | 4 | 填写/修改病例报告表 |
| 5 | 数据疑问解决 | | 6 | | 研究药物接收/清点 | | 7 | 研究药物发放及回收 | | 8 | 研究用品物资管理 |
| 9 | 血液样本采集 | | 10 | | 特殊样本管理 | | 11 | 报告不良反应事件 | | 12 | 与伦理委员会联络 |
| 13 | 应急信封管理 | | 14 | | 质控管理 | | 15 | 其他 | |  |  |
| 参与项目研究人员信息填写（请根据项目情况自行增添） | | | | | | | | | | | |
| 姓名 | | 科室 | | 职称 | | 职责 | | | 研究分工 | | |
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