内审实施计划

AYJ/QR822-002 A/0

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| 审核目的 |  | | | | |
| 审核范围 |  | | | | |
| 审核性质 |  | | | | |
| 审核依据 |  | | | | |
| 审核时间 |  | | 审核组长 | |  |
| 内审员 |  | | | | |
| 日程安排 |  | | | | |
| 首次会议 |  | | 参加人员 | |  |
| 日期 | 受审部门 | 审核时间 | | 审核要素（按医疗器械生产质量管理规范现场检查指导原则条款自查） | |
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| 末次会议 |  | | 参加人员 | |  |
| 备注 |  | | | | |

编制： 批准：