

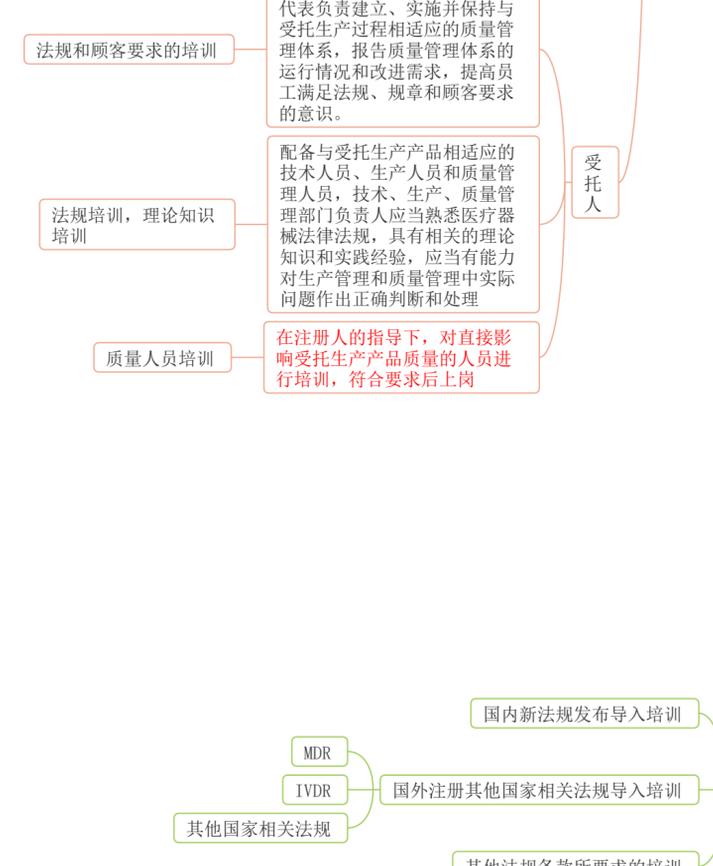
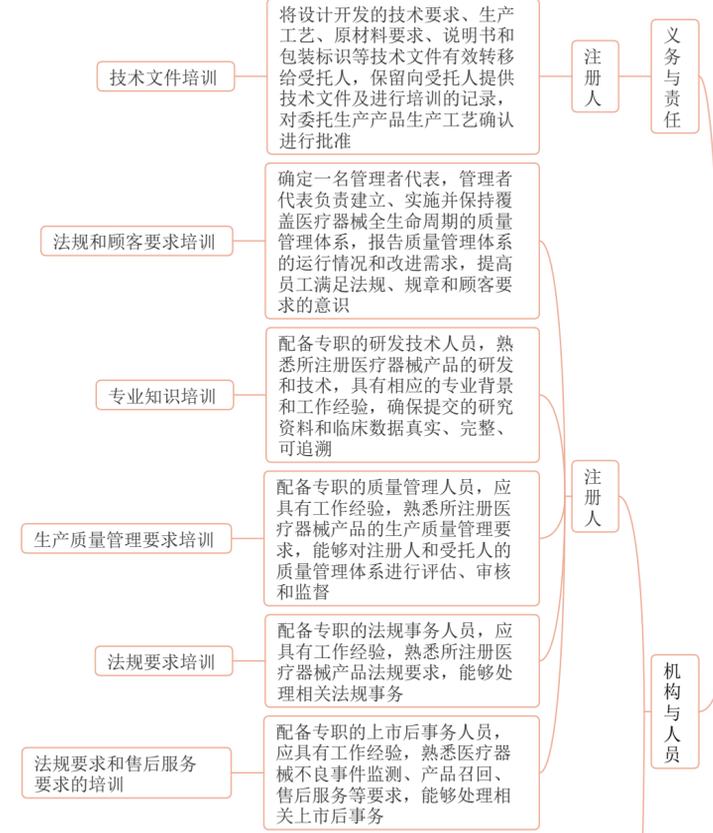
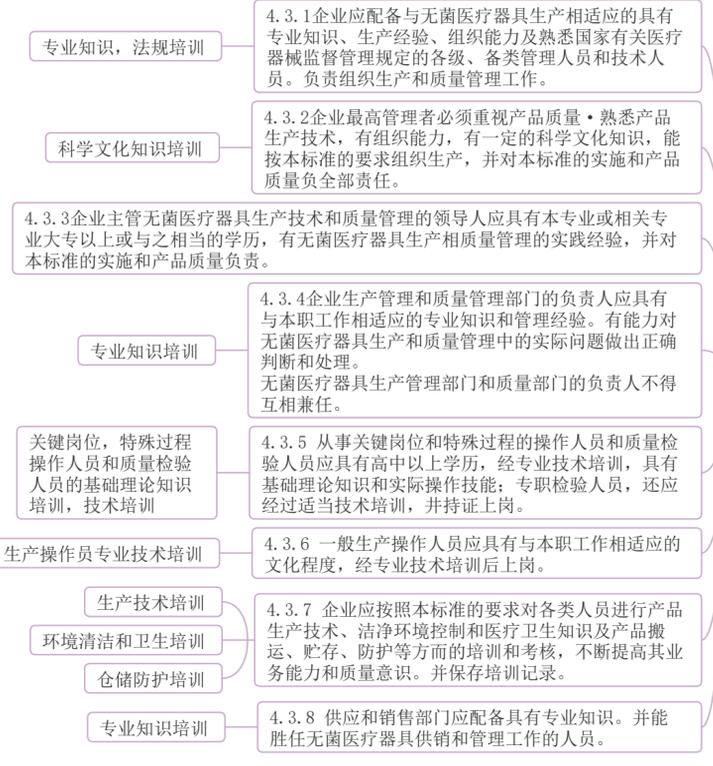
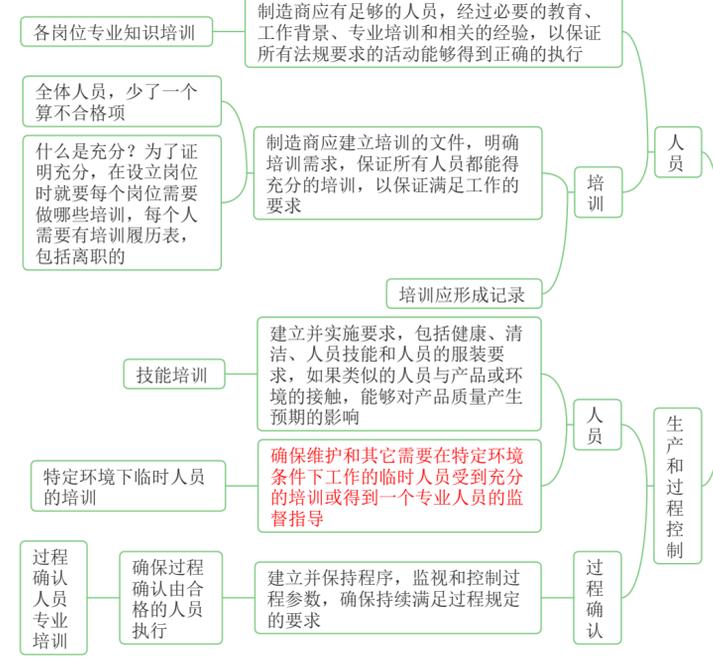
医疗器械企业需要做哪些培训

QSR820

YY0033

注册人制度

新法规导入培训



培训分类

ISO13485

医疗器械生产质量管理规范

医疗器械注册人不良事件监测

