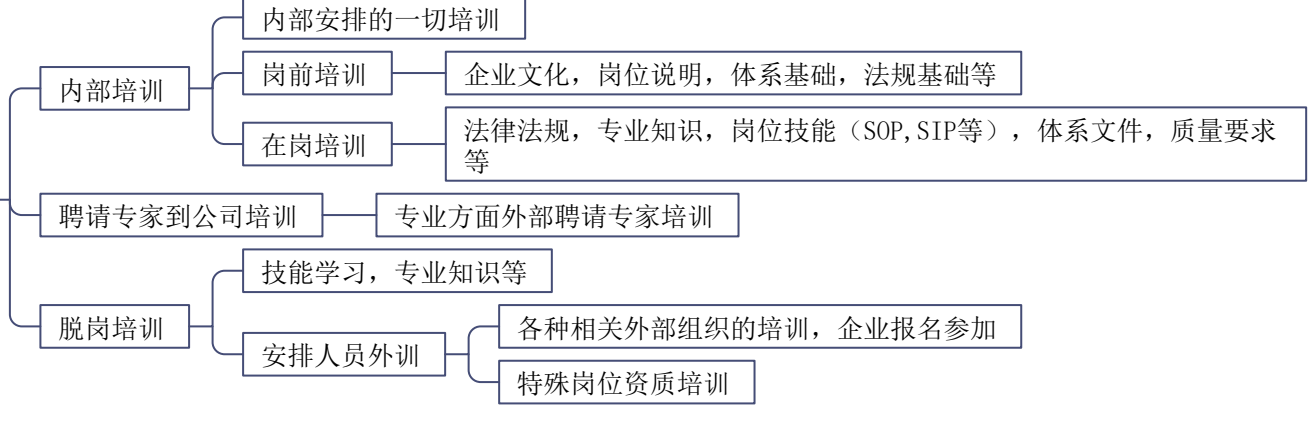
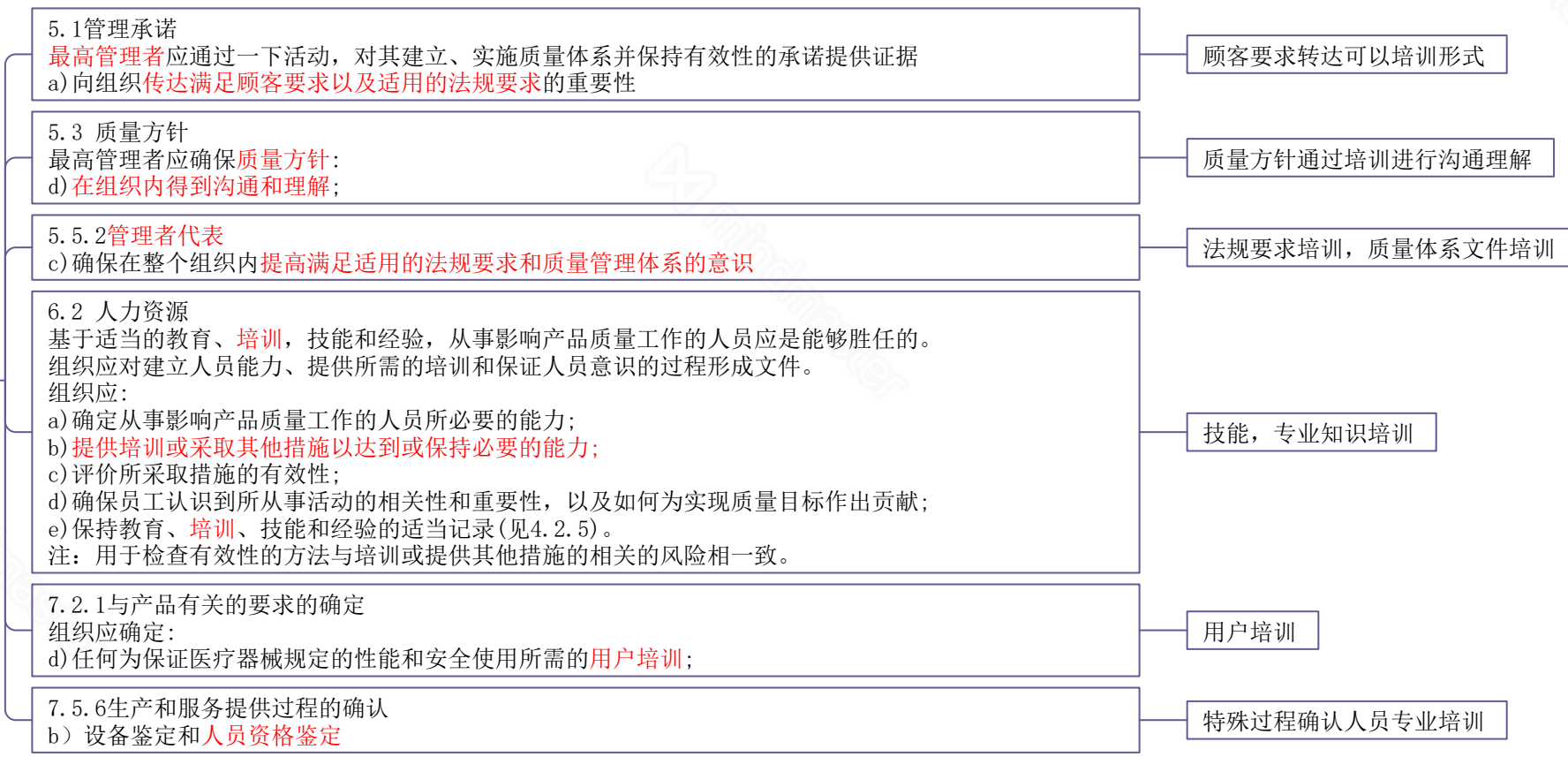


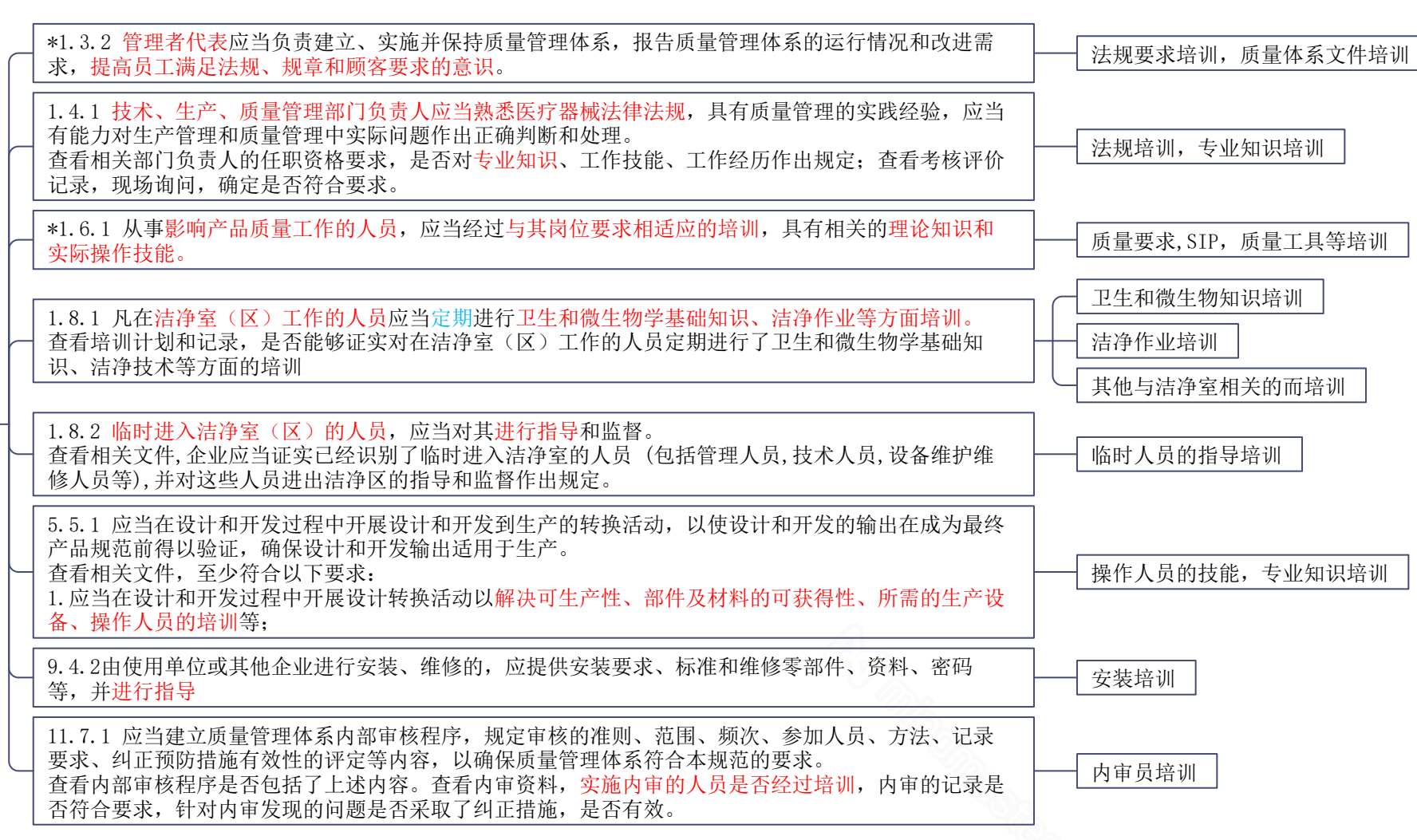
# 培训分类



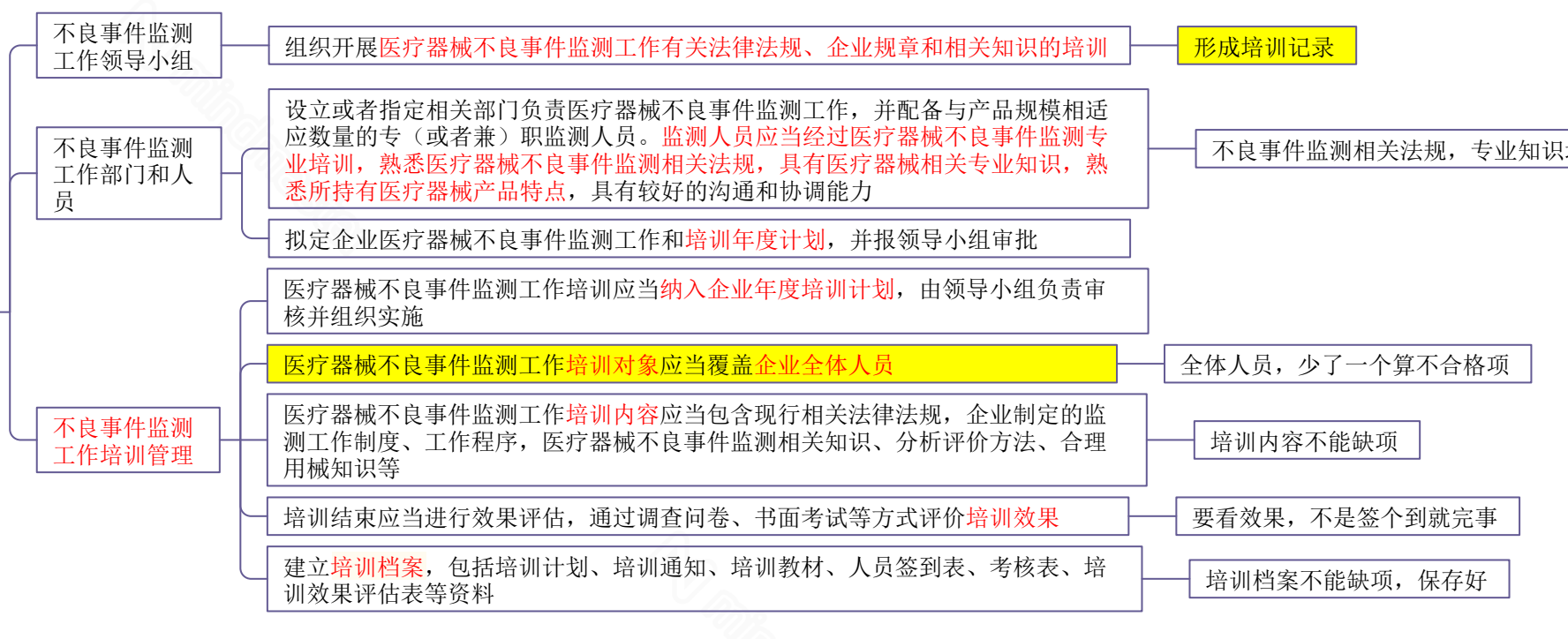
# ISO 13485



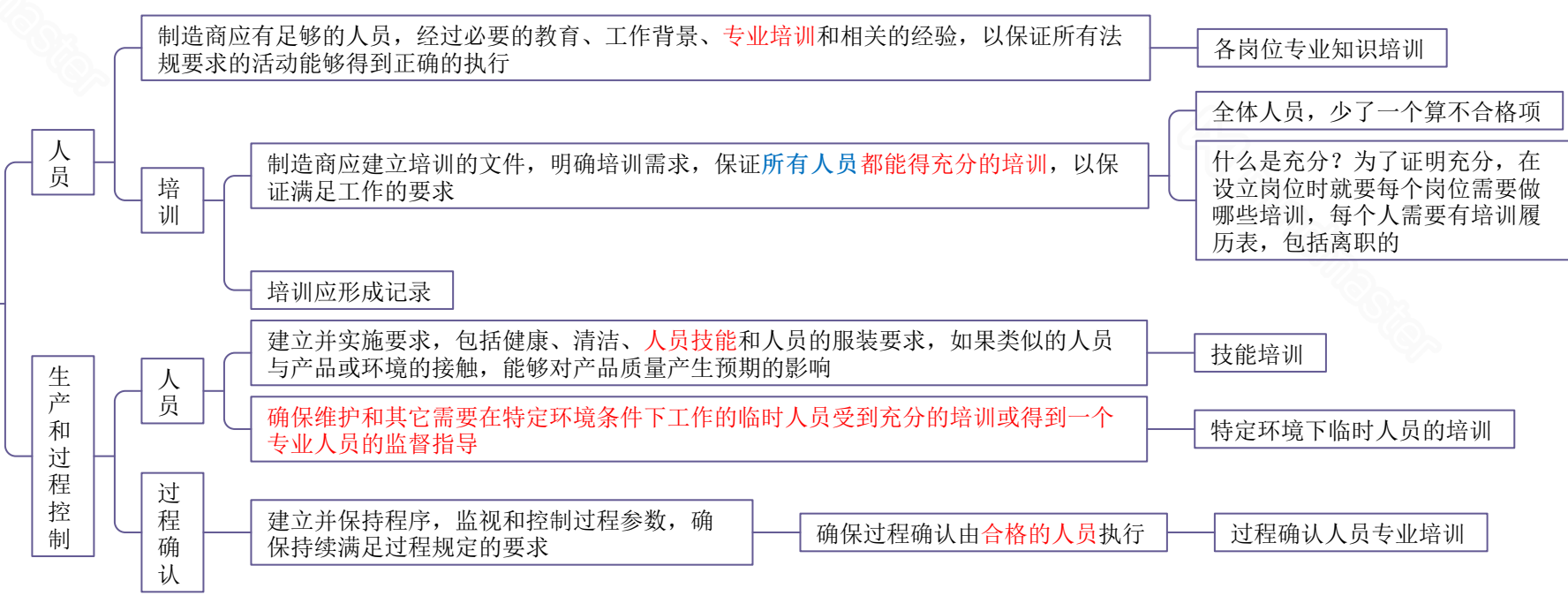
# 医疗器械生产质量管理规范



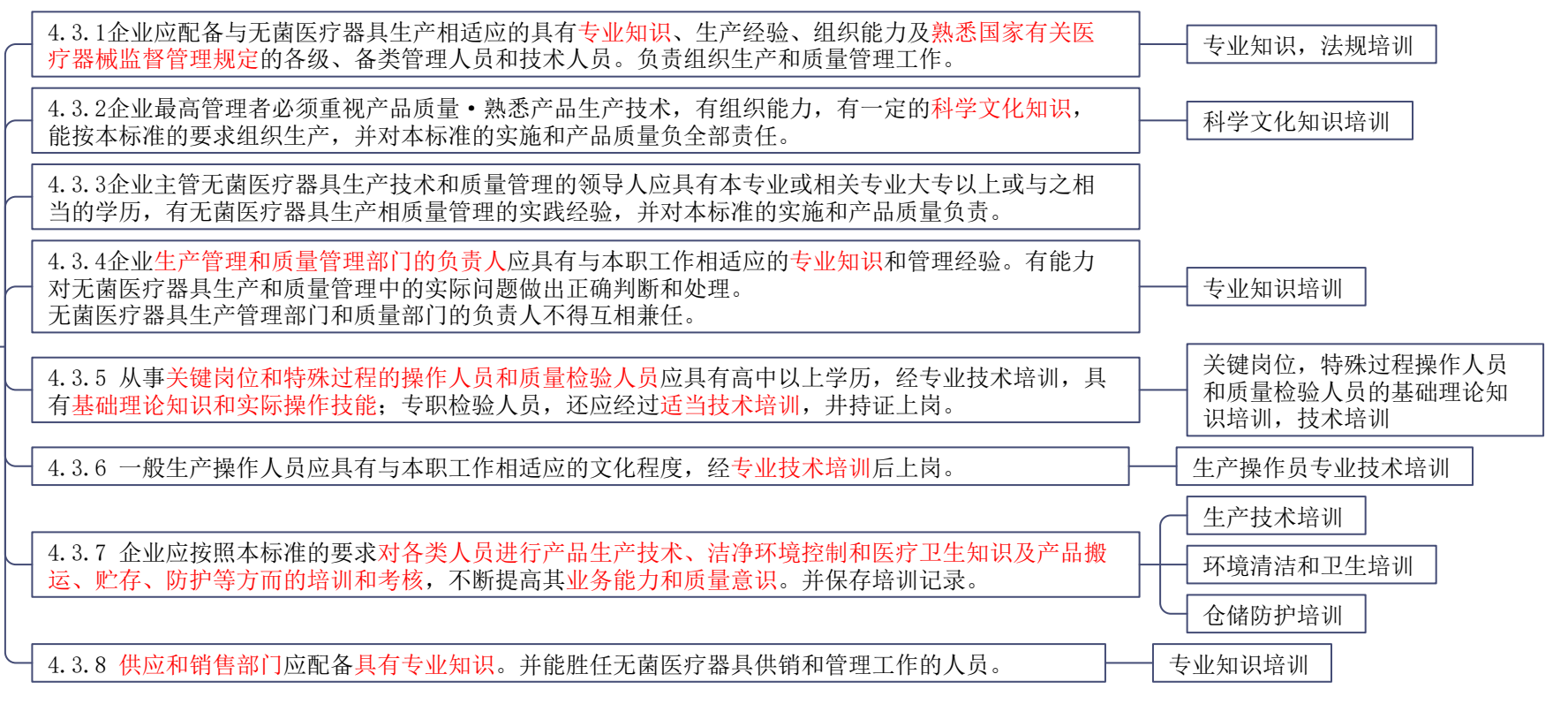
# 医疗器械注册人不良事件监测



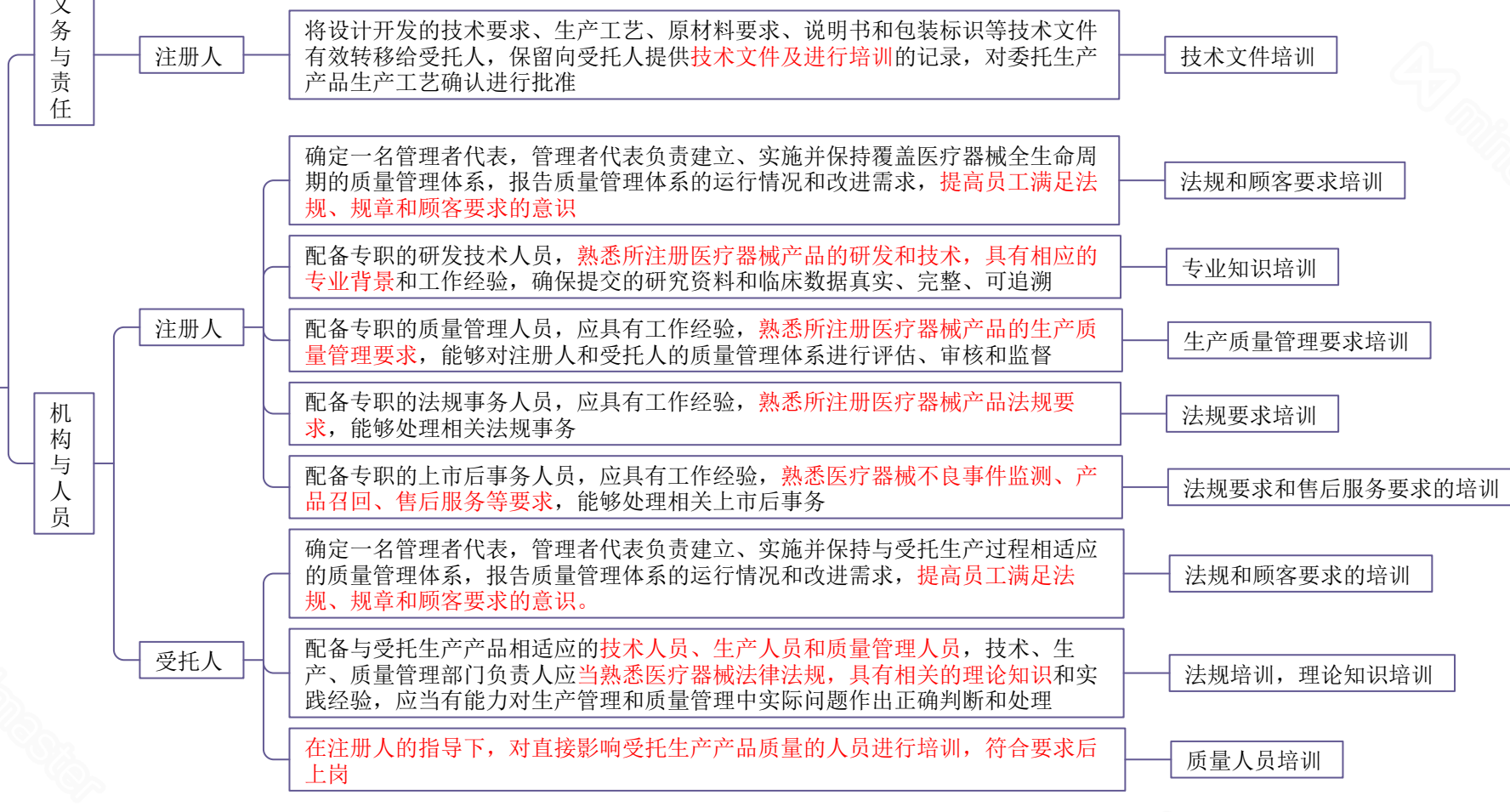
# QSR 820



# YY 0033



# 注册人制度



# 新法规导入培训

