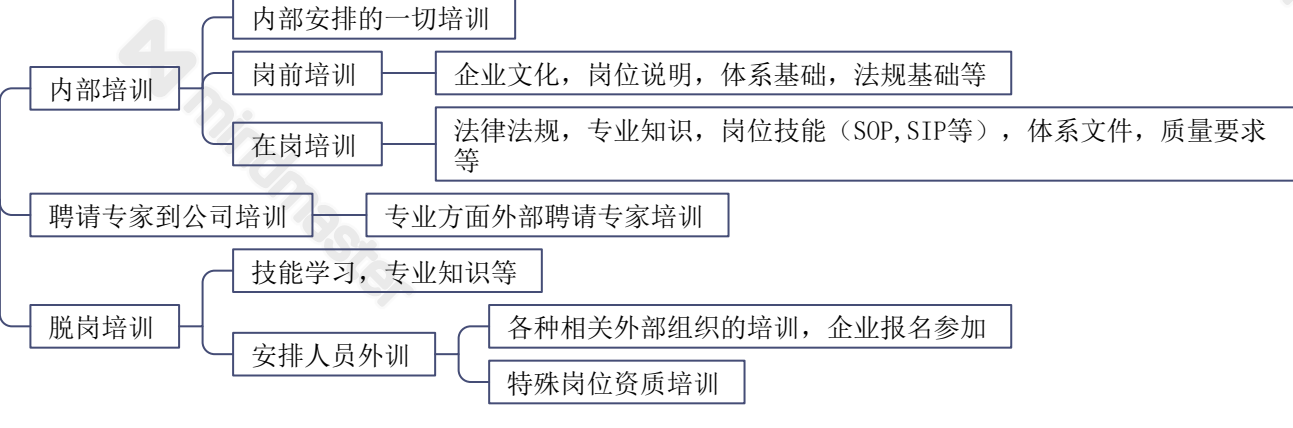
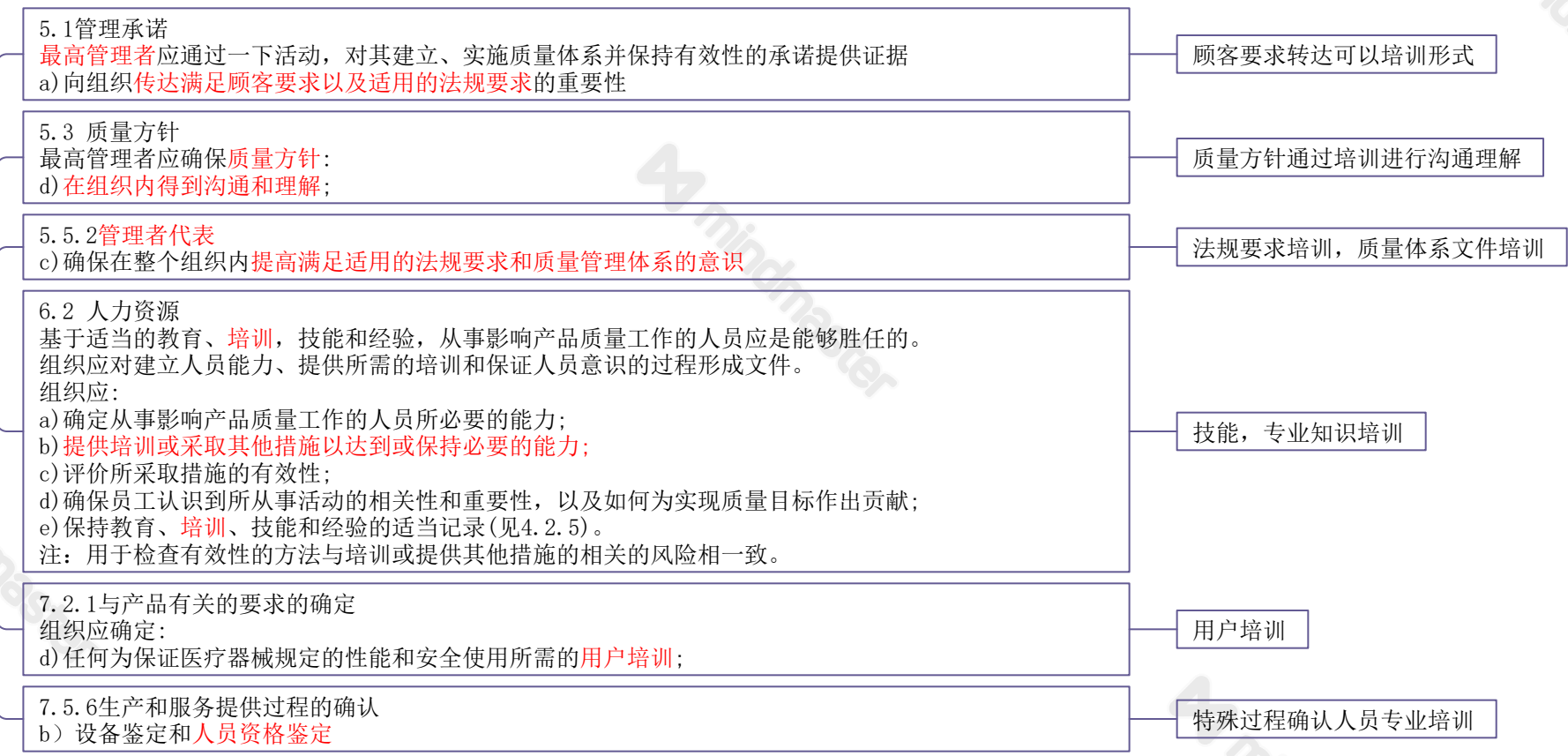


培训分类



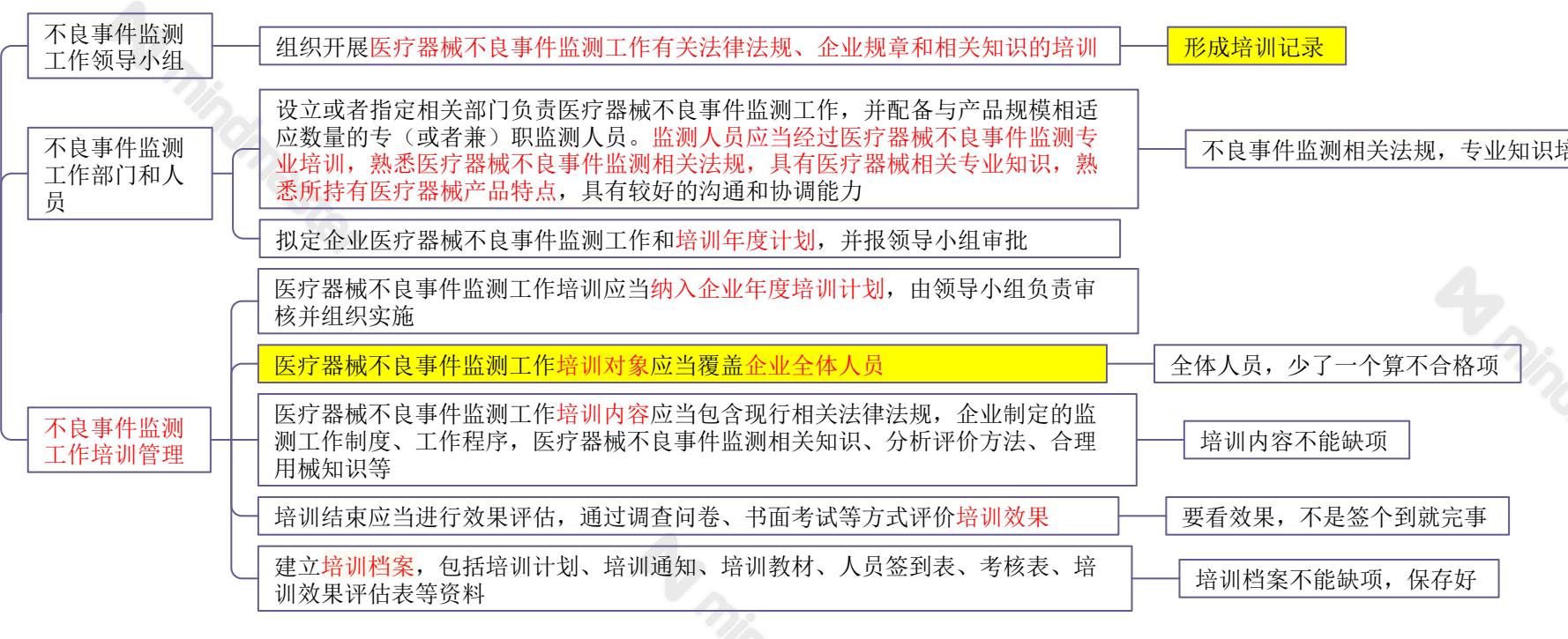
ISO13485



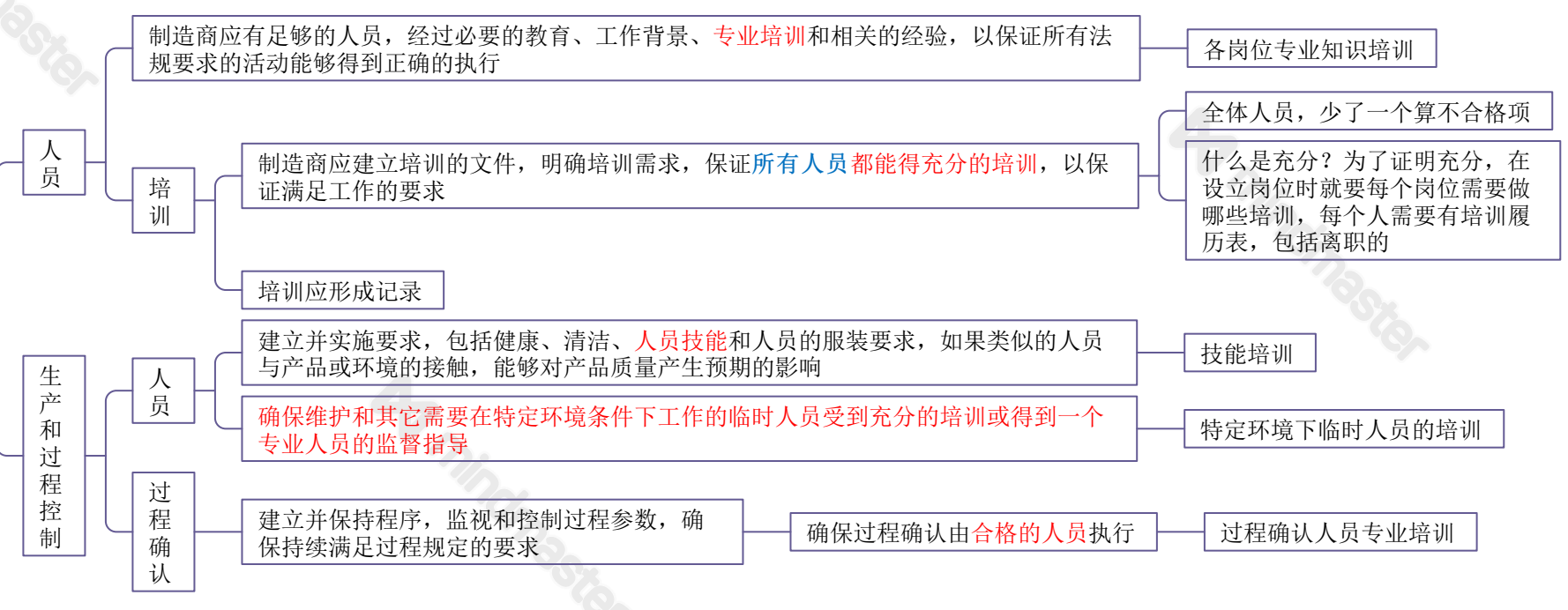
医疗器械生产质量管理规范



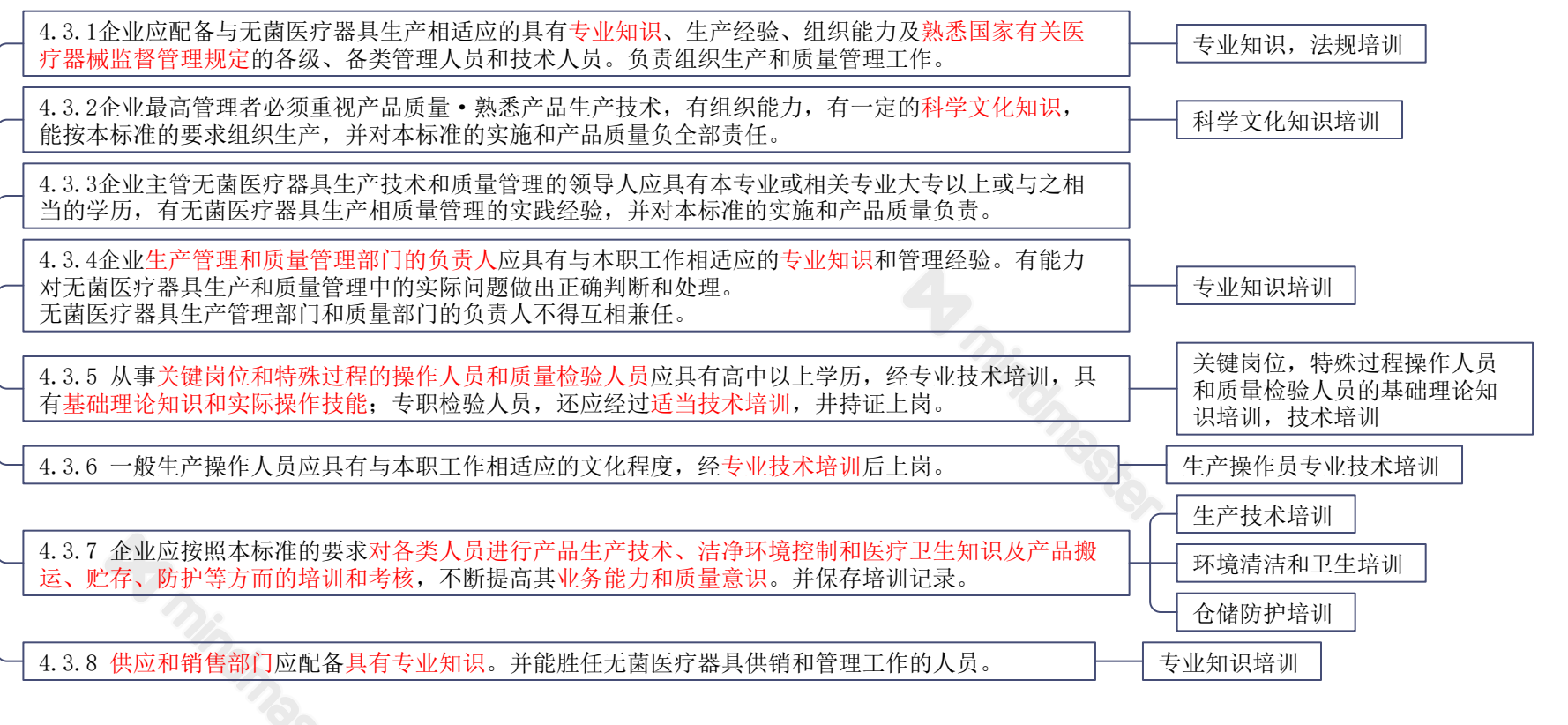
医疗器械注册人不良事件监测



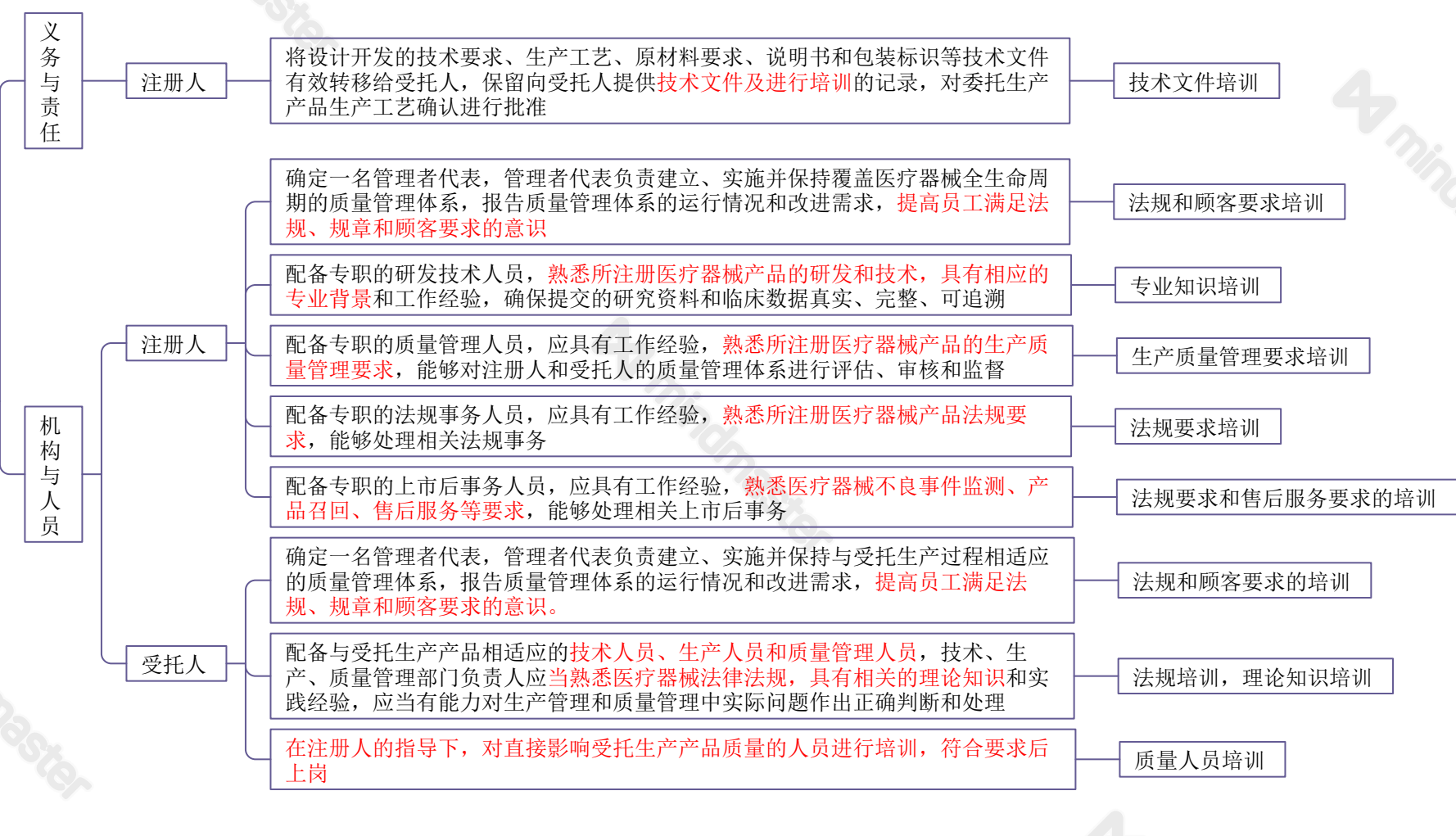
QSR820



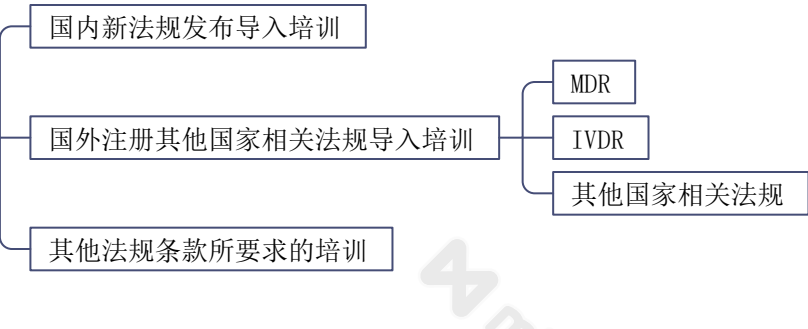
YY0033



注册人制度



新法规导入培训



医疗器械企业需要做哪些培训

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本图是个人理解医疗器械法规整理，用于特定群体实时交流并更新，参照对应法规适用。

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