References & Recommended Reading

1. WHO good manufacturing practices for pharmaceutical products: main principles. In: WHO Expert Committee on Specifications for Pharmaceutical Preparations: forty-eighth report. Geneva: World Health Organization; 2014: Annex 2 (WHO Technical Report Series, No. 986)

2. Supplementary guidelines on good manufacturing practice: validation. In: WHO Expert Committee on Specifications for Pharmaceutical Preparations: fortieth report. Geneva: world Health Organization; 2006: Annex 4 (WHO Technical Report Series, No. 937).

3. Supplementary guidelines on good manufacturing practice: validation. Qualification of systems and equipment. In: WHO Expert Committee on Specifications for Pharmaceutical Preparations: fortieth report. Geneva: World Health Organization; 2006: Annex 4, Appendix 6 (WHO Technical Report Series, No. 937).

4. Supplementary guidelines on good manufacturing practices: validation. Validation of computerized systems. In: WHO Expert Committee on Specifications for Pharmaceutical Preparations: fortieth report. Geneva: World Health Organization; 2006: Annex 4, Appendix 5 (WHO Technical Report Series, No. 937).