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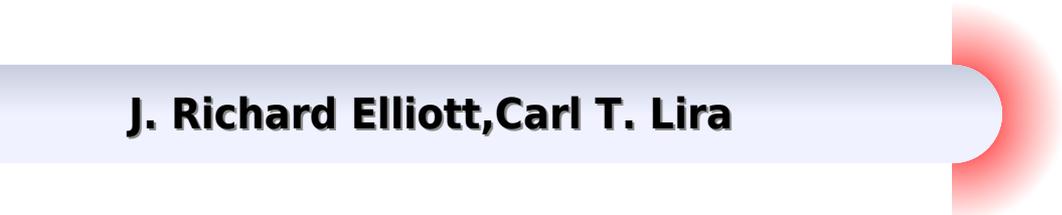
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Analytical Chemistry in a GMP Environment James M. Miller, Jonathan B. Crowther, 2000-05 Based on the Laboratory Analyst Training and Certification Program chemists from a range of pharmaceutical companies and a few academic laboratories explain how to comply with the US Food and Drug Administration s Good Manufacturing Practice rules as analytical technologies are changing rapidly Among the topics are the drug development process uniform and consistent interpretation of compliance issues the role of statistics and basic topics in analytical chemistry and detectors and quantitative analysis The emphasis is on high performance liquid chromatographic methods

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