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economies worldwide are collectively contributing multiple stresses on aquatic ecosystems by the release of numerous contaminants This in turn demands that basic toxicological information on their potential to harm living species be available Hence environmental protection programs aimed at preserving water quality must have access to comprehensive toxicity screening tools and strategies that can be applied reliably and universally While a good number of toxicity testing procedures and hazard assessment approaches have been published in the scientific literature over the past decades many are wanting in that insufficient detail is available for users to be able to fully understand the test method or scheme and to be able to reproduce it successfully Even standardized techniques published in recognized international standard organization documents are often lacking in thoroughness and minutiae Paucity of information relating to biological test methods may be consequent and trigger several phenomena including generation of invalid data and resulting toxicity measurements erroneous interpretation and decision taking with regards to a particular chemical or environmental issue or simply abandonment of testing procedures Clearly improperly documented toxicity testing methods can be detrimental to their promotion and use as they open the doorway to unnecessary debate and criticism as to their raison d tre Furthermore this situation can indirectly contribute to delaying minimizing or eliminating their application thereby curtailing the important role toxicity testing plays in the overall protection and conservation of aquatic ecosystems Validation of **Chromatography Data Systems** Robert D. McDowall, 2016-11-25 Guiding chromatographers working in regulated industries and helping them to validate their chromatography data systems to meet data integrity business and regulatory needs This book is a detailed look at the life cycle and documented evidence required to ensure a system is fit for purpose throughout the lifecycle Initially providing the regulatory data integrity and system life cycle requirements for computerised system validation the book then develops into a guide on planning specifying managing risk configuring and testing a chromatography data system before release This is followed by operational aspects such as training integration and IT support and finally retirement All areas are discussed in detail with case studies and practical examples provided as appropriate The book has been carefully written and is right up to date including recently released FDA data integrity quidance It provides detailed quidance on good practice and expands on the first edition making it an invaluable addition to a chromatographer s book shelf

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