Guidance on Writing a Quality Assurance Manual

Excerpt from, "Requirements for ASME NQA-1 Certification", Issue No.: 3, Date, February 10, 2014:

- 3.1 The Quality Assurance Manual shall not be a reiteration of the NQA-1 Standard, but rather, a detailed written document which clearly describes the controls for each element of the Nuclear Quality Assurance Program as to who, what, when, where and how the program is being implemented. The Quality Assurance Manual need not be in the same format or sequential arrangement as in the NQA-1 Standard, as long as all the applicable requirements have been covered. The Manual accepted by ASME is the guide for performing the audits and determining continued compliance with the accepted Quality Assurance Program.
 - 3.1.1 It is not a requirement nor desirable that detailed written technical procedures, work instructions, travelers, and/or forms be a part of the Quality Assurance Manual, but rather, the manual satisfactorily address the controls of the applicable NQA-1 Standard requirements for documents that specify quality requirements or prescribe activities affecting quality.

The following is to clarify paragraph 3.1 & subparagraph 3.1.1, and to provide guidance on the type of Quality Assurance (QA) Manuals ASME Audit Teams will be requiring Applicants to provide.

The purpose of the QA Manual is to provide a detailed written document that describes the QA Program with commitment statements identifying which Requirements of Part I and Part II of the NQA-1 Standard are applicable and which are not applicable. The QA Manual will also:

- 1) describe the organization,
- contain senior management's commitment to establishing and maintaining an adequate and effective QA Program, and
- 3) identify the product (items/services) provided under the QA Program, and
- describe the processes used to supply the product (items/services).

The QA Manual is to clearly identify the product (items/services) being provided under the QA Program and provides an understanding of the processes affecting quality which are implemented under the QA Program to supply the product. To understand the processes, specific activities need to be identified as well as the technique used to implement the process/activity. The processes affecting quality are those elements which are addressed by a Requirement (NQA-1, Part I) or Subpart II work practice (NQA-1, Part II) that is relevant in attaining and verifying quality in the supply of the product. The technique used to implement these processes are the "controls" (who, what, when, where and how) which will be audited by ASME. Completing the QA Manual Checklist will serve as an aid in determining whether the QA Manual is an auditable working document with sufficient and adequate "controls" to retain its functionality as an ASME audited working document while permitting future revisions to sub-tier documents under the QA Program to be made in accordance with the controls specified in the QA Manual without need for additional ASME audits. These sub-tier documents are the drawings, procedures, work instructions, forms, travelers and/or tags used to convey specific detailed instructions/information to either complete or record a specific activity. The QA Manual is to prescribe a process which provides a high level of

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Table of Contents Guidance Writing Quality Manual

- 1. Understanding the eBook Guidance Writing Quality Manual
 - The Rise of Digital Reading Guidance Writing Quality Manual
 - Advantages of eBooks Over Traditional Books
- 2. Identifying Guidance Writing Quality Manual
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
- 3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Guidance Writing Quality Manual
 - User-Friendly Interface
- 4. Exploring eBook Recommendations from Guidance Writing Quality Manual
 - Personalized Recommendations
 - Guidance Writing Quality Manual User Reviews and Ratings
 - Guidance Writing Quality Manual and Bestseller Lists

- 5. Accessing Guidance Writing Quality Manual Free and Paid eBooks
 - Guidance Writing Quality Manual Public Domain eBooks
 - Guidance Writing Quality Manual eBook Subscription Services
 - Guidance Writing Quality Manual Budget-Friendly Options
- 6. Navigating Guidance Writing Quality Manual eBook Formats
 - o ePub, PDF, MOBI, and More
 - Guidance Writing Quality Manual Compatibility with Devices
 - Guidance Writing Quality Manual Enhanced eBook Features
- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Guidance Writing Quality Manual
 - Highlighting and Note-Taking Guidance Writing Quality Manual
 - Interactive Elements Guidance Writing Quality Manual
- 8. Staying Engaged with Guidance Writing Quality Manual
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Guidance Writing Quality Manual
- 9. Balancing eBooks and Physical Books Guidance Writing Quality Manual
 - \circ Benefits of a Digital Library
 - Creating a Diverse Reading Collection Guidance Writing Quality Manual
- 10. Overcoming Reading Challenges
 - o Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Guidance Writing Quality Manual
 - Setting Reading Goals Guidance Writing Quality Manual
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Guidance Writing Quality Manual
 - Fact-Checking eBook Content of Guidance Writing Quality Manual
 - Distinguishing Credible Sources
- 13. Promoting Lifelong Learning

- Utilizing eBooks for Skill Development
- Exploring Educational eBooks
- 14. Embracing eBook Trends
 - Integration of Multimedia Elements
 - Interactive and Gamified eBooks

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