

NEXT Laboratory Quality Assurance Program Description

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Meeting Focus



- Basis:
 - Discuss the Basis of the ACU NEXT Lab Quality Assurance Program
- Scope:
 - Describe the ACU NEXT Lab Quality Assurance Scope
- Submission:
 - Discuss ACU's proposal to submit the Quality Assurance Program for approval before submission of the Construction Permit (CP) application.



SOME DEFINITIONS

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• QUALITY

• The degree to which an item or process meets or exceeds requirements and expectations.

• QUALITY ASSURANCE

• Those planned and systematic actions necessary to provide adequate confidence that the structure, system or component will perform satisfactorily in service.

SAFETY-RELATED STRUCTURES, SYSTEMS AND COMPONENTS

- Those Structures, Systems and Components (SSCs) that are relied upon to remain functional during and following design basis events to assure:
- 1) The integrity of the fission product physical boundary;
- 2) The capability to shut down the reactor and maintain it in a safe shutdown condition; or
- 3) The capability to prevent or mitigate the consequences of accidents which could result in potential offsite exposures comparable to the applicable guideline exposures set forth in 10 CFR Part 20.



Quality Assurance Basis



10 CFR 50.34(a)(7) requires each applicant for a construction permit to build a production or utilization facility to include, in its preliminary safety analysis report, a description of the quality assurance program to be applied to the design and construction of the structures, systems, and components of the facility

NRC Regulatory Guide 2.5 – "Quality Assurance Program Requirements for Research and Test Reactors"

• Endorses ANSI/ANS 15.8

The general requirements for establishing and executing a quality assurance program for the design, construction, testing, modification, and maintenance of research and test reactors in ANSI/ANS-15.8-1995 provide an acceptable method for complying with the program requirements of 10 CFR 50.34, "Contents of Applications; Technical Information."



Quality Assurance Scope

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The Quality Assurance Program is implemented using a Graded Approach

Designate safety classification by the safety significance of the reactor design.

3 Safety Class, or QA Categories:

- 1. Safety-related (SR) SSCs, as defined earlier, means those structures, systems and components that are relied upon to remain functional during and following design basis events.
- 2. Augmented Quality SSCs support Safety-Related SSCs and have been designated as requiring additional quality level oversight.

This designation may be applied to any item that is subject to non-safety related regulatory requirements but may benefit from additional rigor.

3. Non-Safety Related (NSR) SSCs are SSCs not considered safety related or important to safety.



Quality Assurance Manual Purpose

The QAM specifies:

- QA Policy
- Quality System implementation
- Organizational structure
- Staff roles, responsibilities, and authorities
- Technical activities
- Corrective Action Program



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Quality System Tools

Planning: QA Program Description (QAPD) Prepared and in internal review

Implementation:

Quality Assurance Manual (QAM)

Prepared and in internal review

Quality Procedures (QPs):

35 identified QPs – Implement QAM Requirements

30 are prepared and in internal review

Work Instructions (WIs)

3 identified WIs – Provide supporting implementing instruction to the QPs

1 prepared and in internal review



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QAM Requirements Cross Walk

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ANSI/ANS 15.8 requirements		QAM Section	
1	Organization	2.1	 Quality assurance hierarchy Organization responsibilities
2	Quality assurance program	2.2	 Quality assurance hierarchy Organization, personnel training and qualification
3	Design Control	2.3 2.5 2.9	 Design control Instructions, procedures, and drawings Control of special processes
4 5	Procurement Document Control Procedures, Instructions and Drawings	2.4 2.5 2.9	 Procurement document control Instructions, procedures, and drawings Control of special processes

QAM Requirements (Cont'd)

ANSI/A	NS 15.8 requirements	QAM section	
6	Document control	2.6	• Document control
7	Control of purchased items and service	s 2.7	 Control of purchased items and services
8	Identification and control of items	2.8	 Identification and control of items

9Control of special processes2.9• Control of special processes10Inspections2.10• Inspections

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QAM Requirements (Cont'd)

Control of nonconforming items and

15

services

ANSI/ANS 15.8 requirements QAM Section 11 Test control 2.11 Test control 12 **Control of measuring and test** 2.12 • Control of measuring and test equipment equipment 13 Handling, storage, and shipping 2.13 • Handling, storage, and shipping 14 Inspection, test, and operating status 2.10 Inspections

2.11

2.14

2.15

- Test control
- Inspection, test, and operating status
- Identification and control of nonconforming items

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QAM Requirements (Cont'd)

ANSI/ANS 15.8 requirements

QAM Section

- 16Corrective actions2.16• Corrective action
- 17Quality Records2.17• Quality assurance records

- 18Assessments2.18
- 19Experimental Equipment2.19

- Audits, assessments, and surveillances
- experimental equipment



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Status of QAM, QPs and WIs

Quality Assurance Manual QAM

Prepared

QP-00-01	Quality Assurance Procedures	Prepared
QP-01-01	Organization	
QP-01-02	Contract Review	Prepared
QP-02-01	Annual Management Review	Prepared
QP-02-02	Quality Orientation and Training	Prepared
QP-02-03	Qualification of Lead Auditor	Prepared
QP-02-04	Inspection and Test Personnel	Prepared
QP-03-01	Design Control	Prepared
QP-03-02	Hardware Requirements Specification	Prepared
QP-03-03	Software Quality Assurance Program	Prepared
QP-03-04	Software Quality Assurance Plans	Prepared
QP-03-05	Software Classification	Prepared
QP-03-06	Software Verification and Validation	Prepared
QP-03-07	Cyber Security	
OP-04-01	Procurement Document Control	Prepared

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Prepared



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Status of QAM, QPs and WIs (cont.) NE XZK

QP-05-01	Work Instructions	Nuclear Energy eXperimental Testing
QP-05-02	Drawing Control	Prepared
QP-06-01	Document Control	Prepared
QP-07-01	Control of Supplier Document	Prepared
QP-07-02	Vendor Evaluation and Approval	Prepared
QP-07-03	Acceptance of Items and Services	Prepared
QP-07-04	Supplier Nonconformances	Prepared
QP-07-05	Commercial Grade Dedication	Prepared
QP-08-01	Identification and Control of Materials, Parts, and Components	Prepared
QP-09-01	Control of Special Processes	
QP-11-01	Test Control	
QP-12-01	Control of Measuring and Test Equipment	Prepared
QP-13-01	Handling Storage and Shipping	Prepared
QP-14-01	Inspection, Test, and Operating Status	Prepared
QP-15-01	Control of Nonconformances	Prepared
QP-15-02	Reporting Defects and Noncompliance to 10CFR Part21	Prepared
QP-16-01	Corrective Action Request	Prepared
QP-17-01	Quality Assurance Records	Prepared
QP-18-01	Quality Assurance Audits	Prepared Ar
QP-19-01	Experiment and Test Control	Ð



Status of QAM, QPs and WIs (cont.) NEXT

WI-03-01	Control of Engineering Calcu	ulations
WI-03-02	Control of Design Analysis	
WI-06-01	Control of Engineering Draw	vings

Prepared



Quality Assurance Submission

ACU purposes:



1. Submit the QA Program in early 2021 for NRC review and approval before submission of the CP application.

2. Pre-CP submittal activities under the QA program:

- Conduct of Research
- Technical Assessments
- R&D Validation and Verification (V&V)
- Construction of a potential future reactor room
- Molten Salt Research Reactor Conceptual Design Work





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