



Franklin Research Center  
A Division of The Franklin Institute

June 27, 1983

Mr. Uldis Potapovs, Chief  
Vendor Program Branch  
Nuclear Regulatory Commission  
Region IV  
611 Ryan Plaza Drive, Suite 1000  
Arlington, Texas 76011

JUL - 5 1983

Reference: Docket No. 99900921/83-01; "Notice of Nonconformance" transmitted as Appendix B in your letter dated June 1, 1983.

Dear Mr. Potapovs:

In response to the findings cited in the referenced Notice of Nonconformance, the Franklin Research Center (FRC) proposes to complete the following corrective actions and preventive measures on or before September 1, 1983.

"Notice of  
Nonconformance"  
Paragraph

Proposed Corrective Actions  
and Preventive Measures

(2) QASD 4-1 A.1. Finished  
(3) QAP 4-1 3 Procedures  
QAP 4-2 ok 10-27-83 (GH)

Detailed documented procedures will be established for:  
(1) the training of technical personnel, (2) quality assurance indoctrination of personnel, and (3) the training of auditors and certification of lead auditor. These procedures will describe how, where, and what training records will be maintained. *Need to Identify who is responsible in procedures also. Review when available.*

QAP 4-2 A.2. being  
QASD 17-1 10-27-83 written  
GH ok 10-27-83

A written procedure will be established to identify, control, and evaluate deviations from approved test procedures. *Should Identify who does what in procedures. Review when available.*

QASD 14-1 A.3-1  
GH ok 10-27-83

The responsibilities of Quality Assurance personnel during qualification testing will be described in a documented procedure to assure compliance of the test program with sponsor technical specifications and quality assurance requirements. *Review when available*

Technical and editorial review of Test Procedures (TPs) for Franklin Institute Laboratory, Inc. (FIRL) Project No. 5653 has been accomplished in accordance with paragraph 2.2 of Franklin Quality Assurance Procedure (QAP) No. 14-1. To prevent recurrence, reviews of TPs will be conducted by competent personnel in a more timely manner. *How will this review be documented & what has been changed to assure competent personnel will do the review. Any procedural changes yes*

QAP 9-1 B. Per Andy  
10-27-83 this should  
GH on 8-1-83 be ok. Review  
ok diversity

In accordance with paragraph C.2 of Section 16 of the Franklin QA Manual, QAP No. 16-1 "Storage, Handling, and Shipping of Test Specimens" has been prepared for items undergoing qualification testing at the Franklin Research Center.

*Need to review.*

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June 27, 1983

"Notice of  
Nonconformance"  
Paragraph

Proposed Corrective Actions  
and Preventive Measures

D., E., F.

The technical deficiencies cited in FIRC Project No. 5653 have been communicated to the sponsor (see D. Paulson's letter to PECO dated 6-9-83, and W. Clune's letter to FRC dated 6-15-83). To prevent recurrence, the Project Manager shall assure the completeness of engineering analysis documents, and have the concurrence of the Department Director, or his designee, concerning the technical competence of individuals assigned to verify the adequacy of engineering analyses. The Quality Assurance Manager shall ensure that the test plans and procedures prepared by the Project Manager incorporate and satisfactorily implement contractual requirements, and are reviewed for approval by qualified personnel. *Is there a change to procedures on Manual to require the above? yes*

A documented procedure will be established which shall delineate the responsibilities of Quality Assurance personnel in monitoring qualification testing programs. A checklist with sign-off by QA personnel will be developed to assure the qualification of test personnel, the identification and control of test specimens, the use of approved test procedures, adequate environmental conditions, and the use of calibrated test instrumentation.

*Review when available*

In addition to the above corrective measures, FRC will clarify the Franklin QA Manual and supplementing procedures as follows:

1. The corporate QA policy statement in Section 2 of the QA Manual will be strengthened. *verify*
2. Section 14. "Test Control" and supporting QAPs will be modified to specifically state that the Quality Assurance Manager, or his designee, will review Test Procedures for the inclusion of QA contractual requirements. *verify*
3. Test Procedures will be controlled under appropriate document control procedures. *verify*

Please let me know if the above actions are an acceptable resolution of your Notice of Nonconformance.

Sincerely yours,

*A. J. Saggiomo*

A. J. Saggiomo  
Quality Assurance Manager

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Attachments