



CBI Services, Inc.

St. George Road
Bourbonnais, Illinois 60914-9799

815 933 2200

10CFR71 QA PROGRAM

FOR

KANKAKEE MANUFACTURING RADIOGRAPHY USING SEALED SOURCES

NRC LICENSE NUMBER 12-05639-01

1. Organization:

The final responsibility for the Quality Assurance Program for Part 71 Requirements rests with CBI Services, Inc. Design and fabrication of radioactive material shipping packages shall not be conducted under this Quality Assurance Program. The Quality Assurance Program is implemented using the organization shown in Figure 1.

The Radiation Safety Officer or Assistant Radiation Safety Officer is responsible for overall administration of the program, training and certification, document control, and auditing.

The Radiographers are responsible for handling, storing, shipping, inspection, test, operating status and record keeping.

2. Quality Assurance Program:

The management of CBI Services, Inc. establishes and implements this Quality Assurance Program. Training for all QA functions, prior to engagement in these functions, is required according to written instructions. QA Program revisions will be made with management approval.

The Radiation Safety Officer or Assistant Radiation Safety Officer shall assure that all radioactive material shipping packages are designed and manufactured under a Quality Assurance Program approved by the Nuclear Regulatory Commission for all packages designed or fabricated after July 1, 1978. This requirement can be satisfied by receiving a certification to this effect from the manufacturer.

3. Document Control:

All documents related to a specific shipping package will be controlled through the use of instructions in the Safety Manual, and all document changes will be approved by management. The Radiation Safety Officer shall insure that all QA functions are conducted in accordance with the latest applicable documents.

4. Handling Storage and Shipping:

Written safety instructions concerning the handling, storage and shipping of packages for certain special form radioactive material will be followed.



4. Handling Storage and Shipping: (Cont.)

Shipments will not be made unless all tests, certifications, acceptances, and final inspections have been completed according to instructions in the Radiation Safety manual.

Radiography personnel shall perform the critical handling, storage and shipping operations.

5. Inspection, Test and Operating Status:

Inspection, test and operating status of packages for certain special form radioactive material will be indicated and controlled by the Radiation Safety Manual. Status will be indicated by tag, label, marking or log entry. Status of nonconforming parts or packages will be positively maintained by written instructions in the Radiation Safety Manual.

Radiography personnel shall perform the regulatory required inspections and tests in accordance with written instructions. The Radiation Safety Officer shall ensure that these functions are performed.

6. Quality Assurance Records:

Records of package approvals (including references and drawings), inspections, tests, operating logs, audit results, personnel training and qualifications and records of shipments will be maintained.

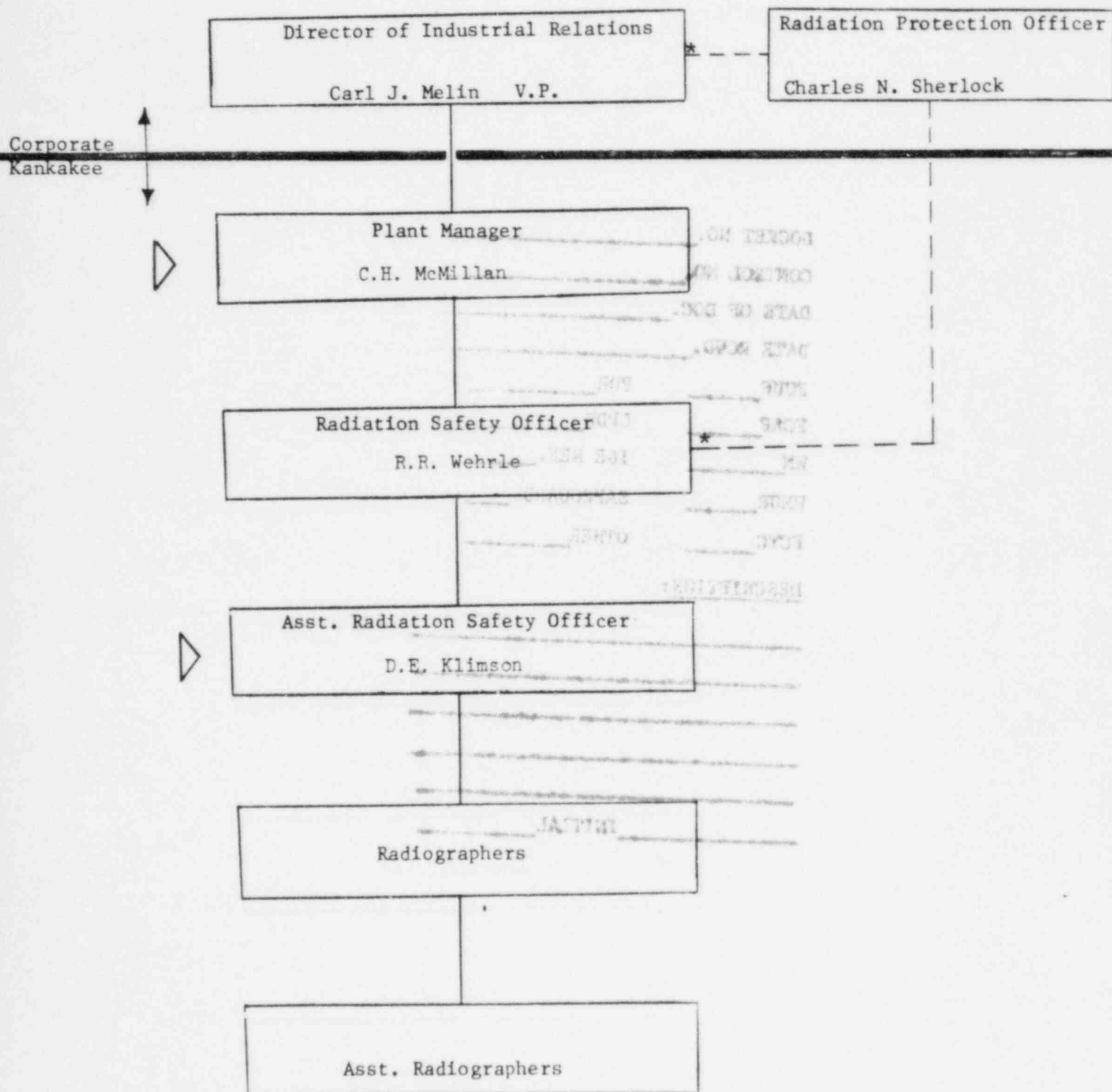
These records will be maintained in accordance with written instructions. The records will be identifiable and retrievable. A list of these records, with their storage locations, will be maintained by the Radiation Safety Officer.

7. Audits:

Established schedules of audits of the Quality Assurance Program will be performed using written check lists. Results of audits will be maintained and reported to management. Audit reports will be evaluated and deficient areas corrected. The audits will be dependent on the safety significance of the activity being audited, but each activity will be audited at least once per year. Audit reports will be maintained as part of the Quality Assurance records. Members of the audit team shall have no responsibility in the activity being audited.



ORGANIZATIONAL STRUCTURE



* - - - - - Denotes Technical Authority

Figure 1.

DOCKET NO. 71-0142
CONTROL NO. 26350
DATE OF DOC. 01/16/86
DATE RCVD. 01/23/86
FCUF _____ PDR ☒
FCAF _____ LPDR _____
WM _____ I&E REF. ☒
WMUR _____ SAFEGUARDS _____
FCTC ☒ OTHER _____

DESCRIPTION:

request amend-
ment to their
Quality Assurance
Program

01/24/86 INITIAL CEC