

TECHNOLOGY PLUS, INC.

10 CFR PART 71 TRANSPORTATION QA PROGRAM
FOR INDUSTRIAL RADIOGRAPHY AND SEALED SOURCES

1. Organization:

The final responsibility for the Quality Assurance (QA) Program for Part 71 Requirements rests with Technology Plus, Incorporated. Design and Fabrication shall not be conducted under this QA Program. The QA Program is implemented using the following organization.

ORGANIZATION CHART TECHNOLOGY PLUS, INC.

PRESIDENT
Garnette Hanson



RSO/MANAGER
Dennis Hanson



RADIOGRAPHERS



ASSISTANT
RADIOGRAPHER

The Radiation Safety Officer is responsible for overall administration of the program, training and certification, document control, and auditing. The Radiation Safety Officer's responsibilities are specifically addressed in Technology Plus, Inc.'s, Radiation Safety Manual, Section 1.

The radiographers are responsible for handling, storing, shipping, inspection, test and operating status and recordkeeping. Radiographers' responsibilities are specifically addressed in Technology Plus, Inc.'s, Radiation Safety Manual, Section 1.

2. Quality Assurance Program:

The management of Technology Plus, Inc., establishes and implements this QA Program. Training, prior to engagement, for all QA functions is required according to written procedures. QA Program revisions will be made according to written procedures with management approval. The QA Program will ensure that all defined QC procedures, engineering

procedures, and specific provisions of the package design approval are satisfied. The QA Program will emphasize control of the characteristics of the package which are critical to safety. Quality Assurance Program specifics are further addressed in Technology Plus, Inc.'s, Radiation Safety Manual, Section 2.

3. Document Control:

All documents relating to a specific shipping package will be controlled through the use of written procedures. All document changes will be performed according to written procedures approved by management.

The Radiation Safety Officer shall insure that all QA functions are conducted in accordance with the latest applicable changes to these documents. Document Control specifics are further addressed in Technology Plus, Inc.'s, Radiation Safety Manual, Section 3, Section 9, and Section 11.

4. Handling, Storage and Shipping:

Written safety procedures concerning the handling, storage, and shipping of packages for radioactive material will be followed. Shipments will not be made unless all tests, certifications, acceptances, and final inspections have been completed. Work instructions will be provided for handling, storage, and shipping operations. Handling, storage and shipping specifics are further addressed in Technology Plus, Inc.'s, Radiation Safety Manual, Section 9.

5. Inspection, Test, and Operating Status:

Inspection, test, and operating status of packages for radioactive material will be indicated and controlled by written procedures. Status will be indicated by tag, label, marking, or log entry. Status of nonconforming parts or packages will be positively maintained by written procedures. Written procedures are specifically addressed in Technology Plus, Inc.'s, Radiation Safety Manual, Section 11 and Section 12.

6. Quality Assurance Records:

Records of package approvals (including references and drawings), procurement, inspections, tests, operating logs, audit results, personnel training and qualifications and records of shipments will be maintained. Descriptions of equipment and written procedures will also be maintained. These records are further addressed in Technology Plus, Inc.'s, Radiation Safety Manual, Section 1, Section 3, Section 9, Section 11, and Section 12.

These records will be maintained in accordance with written procedures. The records will be identified 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 QA 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. Audits are specifically addressed in Technology Plus, Inc.'s, Radiation Safety Manual, Section 13.