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REGISTRANTS TOP TEN VIOLATIONS

1. Not maintaining the appropriate records (Notice of Registration (NOR), Plan Review, Report of Assembly (FDA 2579, Letter of Acknowledgement (LOA), Post room survey, annual review of Written Safety program).
2. The facility Written Radiation Protection program was not available or not adequate.
3. The registrant failed to have a copy of the "North Carolina Regulations For Protection Against Radiation" at the facility.
4. The registrant failed to annually review the Written Radiation Protection program.
5. There has not been an area radiation survey done within 30 days following initial operation of equipment.
6. The registrant failed to provide a working technique chart for each diagnostic x-ray system.
7. The registrant failed to update the information contained in the application for registration or notice of registration no longer accurate.
8. The registrant failed to have the floor plan and equipment arrangement reviewed by a qualified expert prior to construction or structural modification.
9. Appropriate radiation caution signs have not been posted.
10. Personnel monitoring equipment has not been supplied and/or used by all occupationally exposed personnel.