

Radiation Control Program

Violations Summary Report -- 2000 Violations

Inspections Performed by X-Ray Inspectors

From 1/1/2000 Through 12/31/2000

Description of Violation	Number of Violations
Compliance test &/or corrective action not performed	2,626
OS&P not complete/not available; certificate, RPP, or NOV not available	1,768
Time-temp. chart, timer, therm. not available; Wrong safelight filter, wattage, &/or distance	870
Collimator indicators, PBL, light field, &/or centers alignment inadequate	712
Annual protective device test not performed	436
Registration or certification: none, not current, expired	434
Patient exposure at skin entrance (ESE) exceeded limits	356
Mammo QC, physicist eval., or physician review of QC not performed; phantom tech. not patier	328
Technique chart not available	324
Personnel not credentialed; operators, physician, physicist	306
Public dose surveys not performed; Dose to public excessive	258
Timer inaccurate or non-reproducible; Exposure untimed	196
Indicators of x-ray production, multiple tubes, battery charge, AEC operation not available	156
Occupational dose not determined; Current year dose not determined	152
'Notice to Employees' not posted	148
Mammo QC failed &/or phantom failed without ceasing mammograms	130
Other x-ray & other mammo	128
Current personnel dose unknown; PM records not available	122
Fluoro collimation (spot film, input phosphor, Min @ max, &/or stepless) not adequate	108
Mammo anti-scatter grids not available; Compression paddle misaligned	104
Output reproducibility &/or mA linearity inadequate	94

Radiation area caution signs not posted; No machine label	74
Annual fluoroscopic entrance exposure measurements not performed	60
Technique factors not indicated prior to exposure: kVp, mA, mAs, time, AEC	60
Technique chart does not match techniques used clinically	56
Inadequate HVL	52
Mammo image not processed w/in 24 hrs.; Batch processing not performed properly	50
Operator collimation inadequate	26
Mammo QC corrective action not performed	24
Inappropriate PM device; PM device used improperly; Supplier not NVLAP	24
Analytical x-ray operators not qualified	20
Patient viewing system not available or not used; Two-way aural communication not available	20
QA/QC records not available	18
Mammo patient film not provided or not retained	18
Therapy annual calibration not performed	18
X-Ray machine not specifically designed for mammography	14
Exposure switch not the 'deadman' type	10
Protective devices &/or holding devices not available or not used	8
Therapy filter interlock or indicator system inadequate	8
CT dose measurements, spot checks not performed	6
Survey instrument calibration not performed	6
Fluoro minimum SSD inadequate	6
Code not used	6
Overexposure report to Agency or individual not performed	4
Code not used	4
Fluoro energized w/o primary barrier in locked position	4
Fluoro EER exceeds limit	4

Deliberate exposure to radiation	2
X-ray machine non-compliant after installation	2
Industrial x-ray interlock bypass warning not posted	2
RSO quarterly audit records not available	2
Mammo AEC not providing constant density; Mammo kVp inaccurate	2
Service company servicing non-registered x-ray machine	2
Code not used	2
Code not used	2
Minimum source to skin distance (SSD) exceeded (radiographic & dental)	2
Diagnostic tube housing inadequate	2
Tube housing assembly not stable or hand held during radiography	2

Report Compiled on: August 09, 2005