MEDICAL RADIOGRAPHIC REPORT FORM

(A FACILITY form is required with this form)

Facility _____ Date Surveyed_____

Registration number _____-_____ Room number _____ Control number_____ Inspector _____

Address (multiple facilities)_____ Person Interviewed_____

Tubehead Manufacturer and serial no._____

Control panel Manufacturer and serial no._____

	Y	N	N/A	DNT	Y (yes), N(no), N/A(not applicable), DNT (did not test) (All reg references preceded by 0400-20-)
1.	○	○	○		Switch at control panel and any remote switch labeled "Caution Radiation". (05-.111(11))
2.	○	○	○		Proper technique chart posted and used. (04-.11(1)(e), 06-.04(4))
3.	○	○	○		Patient and control panel visible at at all times. (06-.05(2)(c)1.)
4.	○	○	○		Operator remains in protected area during exposures *[for stationary units]*. (06-.05(2)(c)5.(ii)(II)I)
5.	○	○	○		Operator 2 meters away from primary beam *[for mobile units]*. (06-.05(2)(c)5.(ii)(II)II))
6.	○	○	○		Battery indicator shows when sufficient charge for operation *[for mobile units]*. (06-.05(2)(a)9.)
7.	○	○	○	○	Multiple tube indicator on control panel and near tube housing. (06-.05(2)(a)10.)
8.	○	○	○		Beam axis indicator for fixed units. (06-.05(2)(c)7.)
9.	○	○	○		Tubehead stable. *(Drifts or vibrates)* (06-.05(2)(a)12.)
10.	○	○	○	○	Cones or diaphragms marked with film size and source-to-image receptor distance (SID) to be used. (06-.05(2)(a)13)
11.	○	○	○		Technician uses collimator correctly. (04-.12(1)(a)2, 06-.04(4), and 06-.05(2)(a)1)
12.	○	○	○	○	Timer terminates exposure at a preset limit. (06-.05(2)(c)5.(i))
13.	○	○	○	○	Exposure interrupt for time > 0.5 seconds. (06-.05(2)(c)5.(ii)(I)I)
14.	○	○	○	○	Exposure not possible if timer set on "0" or "OFF". (06-.05(2)(c)5,(i)(II))
15.	○	○	○	○	When the unit is in the automatic exposure control mode (AEC), does the AEC conform to the requirements of 0400-20-06-.05(2)(c)5.(iii)?
16.	○	○	○	○	Primary barrier adequate. (05-.60 and 06-.04(2))

☐ Unit is in storage and not being used.

	P	F	N/A	DNT	P(pass), F(fail), N/A(not applicable), DNT (did not test)
17.	○	○	○	○	Is the system provided with means to align the center of the x-ray field to the center of the image receptor to within 2% of the SID (distance from x-ray field center to crosshair or equivalent)? (06-.05(2)(c)7.)
18.	○	○	○	○	Distance from the x-ray field center to the light field center less than 2% of test SID. (06-.05(2)(c)2.)
19.	○	○	○	○	Field Size Indicators (FSI) accurate to within 2% of test SID. (06-.05(2)(c)2.) *(If question 19 is applicable, then questions 26 and 27 are N/A)*
20.	○	○	○	○	Difference in dimensions of the light field and the x-ray field less than 2% of the test SID. (06-.05(2)(c)2.)
21.	○	○	○	○	Distance from the x-ray field center to the bucky film center less than 2% of the bucky SID. (06-.05(2)(c)4 and 7.)
22.	○	○	○	○	Source-to-Image distance indicator (SIDI) accurate to within 2% of the bucky SID. (06-.05(2)(c)7.)
23.	○	○	○	○	Exposure reproducible. (06-.05(2)(c)5.(iv))
24.	○	○	○	○	Timer reproducible. (06-.05(2)(c)5.(v))
25.	○	○	○	○	Millimeters of Al HVL/Filtration adequate. (06-.05(2)(a)2.(i) or (ii))
26.	○	○	○	○	For radiographic equipment designed for only one image receptor size at a fixed SID, are the dimensions of the x-ray field no greater than the dimensions of the image receptor? (06-.05(2)(c)4.) *(If question 26 is applicable, then questions 19 and 27 are N/A)*
27.	○	○	○	○	When using a collimating technique other than the use of FSI (*question 19*) or light field (*when the film or film cassette is not directly visible – question 20*) or when the equipment is designed for only one image receptor size at a fixed SID (*question 26*), is the x-ray field less than the dimensions of the image receptor by 3% in either length or width and 4% by summing length and width excess? (06-.05(2)(a)1.) *(If question 27 is applicable, then questions 19 and 26 are N/A)*