

Nevada Radiation Control Program

Mammography Machine **Application for Certificate of Authorization**



FACILITY NAME (CERTIFICATE ISSUED IN THIS NAME)	TELEPHONE NUMBER		FAX NUMBER	E-MAIL ADDRESS
MAILING ADDRESS	CITY	STATE	ZIP CODE	
PHYSICAL ADDRESS (IF DIFFERENT)	CITY	STATE	ZIP CODE	
NAME OF INDIVIDUAL COMPLETING FORM ¹	TITLE	TELE	PHONE NUMBER	E-MAIL ADDRESS
TE OF PHYSICIAN WHO SUPERVISES MACHINE OPERATION				

LOCATION OF MAMMOGRAPHY PRACTICE

private radiology office	hospital	multi-sp	oecialty cl	inic	mobile unit, fixed use location		
mobile unit, multiple temporary	locations ²	other					
					SPECIFY LOCATION		
Number of mammography machine	Number of mammography machines in use at the designated location?						
Number of mammograms performed each month at this location?							
Proportion of total mammograms p	erformed for clinic	<u></u> %					
Are non-breast imaging studies per	formed at this loca	ation?	Υ	N			

If not a member of organizational management, provide a letter from management which authorizes this individual to legally bind the organization.
 List use locations on supplemental attached sheet.

³ Non-screening; ordered because of symptoms, findings, or prior history of breast cancer.

PERSONNEL 4

RADIOLOGIST	ABR CERTIFIED	YEAR OF ABR CERT	NEVADA LIC NO.	YEARS EXPERIENCE IN MAMMOGRAPHY	NAMED ON PRIOR CERT	DIGITAL QUALIFIED
	Y N				Y N	Y N
	-					

NAME OF RADIOLOGICAL PHYSICIST FOR FACILITY ⁵	SPECIALTY BOARD CERTIFIED	YEAR OF SPECIALTY CERTIFICATION	SUBSPECIALTY BOARD CERTIFICATION	DIGITAL QUALIFIED
	Y N		Y N	Y N

⁴ Name all physicians interpreting mammograms for this facility. Indicate if they are ABR certified radiologists and any special mammography training (e.g., residency, fellowship, workshops, seminars, symposia, etc.). Provide the number of years of mammography interpretation experience for each.

This physician supervises machine operation for purposes of this application.

NAME OF MAMMOGRAPHY TECHNOLOGIST	AART CERTIFIED C		NEVADA CERTIFICATION NO.	DIGITAL QUALIFIED
	Υ	N		Y N
CHIEF TECHNOLOGIST				
SUPERVISING TECHNOLOGIST ⁶				

TECHNIQUES AND POSITIONS USED FOR NON-SYMPTOMATIC MAMMOGRAPHY								
How many views per breast comprise a typical non-symptomatic exam?								
CC	ML oblique	ML (chest wall)	9	O° ML7 (contact)	other			
						SPECIFY		
Typical technique employed for a craniocaudad view of an average density (normal) compressed breast:								
		Grid used?	Υ	N				
kVp	mA TIME	mAs						

⁶ For purposes of this application, the supervising technologist may or may not hold a Certification of Authorization for Mammography.

EQUIPMENT 7

Previous Health Division Radiation Produc	ng Machine	Registration or Certificate?	YCERTIFICATE NUMBER	N
Dedicated mammography machine?	′ N			
		MANUFACTURER	MODEL NUMBER	GENERATOR SERIAL NUMBER
DATE OF MANUFACTURE DATE INSTALLED	FOCUS	RECEPTOR DISTANCE (cm)		
Target material: W Mo Rh	Ag	other		
Previous Health Division Radiation Produc	ng Machine	Registration or Certificate?	Υ	_ N
Dedicated mammography machine?	' N		CERTIFICATE NUMBER	
3 1 3		MANUFACTURER	MODEL NUMBER	GENERATOR SERIAL NUMBER
DATE OF MANUFACTURE DATE INSTALLED	FOCUS	RECEPTOR DISTANCE (cm)		
Target material: W Mo Rh	Ag	other		
Previous Health Division Radiation Produc	ng Machine	Registration or Certificate?	Υ	_ N
Dedicated mammography machine?	' N		CERTIFICATE NUMBER	
		MANUFACTURER	MODEL NUMBER	GENERATOR SERIAL NUMBER
DATE OF MANUFACTURE DATE INSTALLED	FOCUS	RECEPTOR DISTANCE (cm)		
Target material: W Mo Rh	Ag	other		

⁷ Complete one section for each mammographic machine to be issued a certificate. Include in your application package a phantom image taken with each machine, and a copy of the post-installation physicist report.

IMAGE RECORDING SYSTEM

	FILM/SCREEN 8						
				FILM			
	Agfa-Mamoray	Dupont LDS	Dupont LDT	Dupont Mic	crov Dupo	ont MRF31	
	Dupont SR329	Fuji-MI	Kodak Mini-R	Kodak OM	Koda	k TMM	
	Konica, CM	Other					
	SCREEN						
	DuPont Lo Dose		Dupont LD/2	Kodak Min-	-R (SSS) ⁹	Kodak Min-R Fast (DSS) 10	
	Kodak Min-R Medium	n (SSS)	Kodak Min-R-RF	Konica	, ,	Monarch	
	other	,					
	s the above a shange fr	om system uses	- H from provious corti	Signato 2 V	NI		
ı	s the above a change fr	om system used	a from previous certii	Ficate? Y	N		
	Digital system			IDENTIFY PRINTER SYSTE	EM		
			PROC	ESSOR 11			
	MANUFACTURER			CHEMISTRY TYPE	OPEPRATING TEMP (C/F)	TIME IN DEVELOPER SOLUTION (SEC)	
	Dedicated to mammogra	iphy only?	Y N				
_							
	MANUFACTURER	MODEL	NUMBER	CHEMISTRY TYPE	OPEPRATING TEMP (C/F)	TIME IN DEVELOPER SOLUTION (SEC)	

Ν

Dedicated to mammography only?

⁸ Identify each type. If film and screen do not bear the same brand, provide manufacturer's statement of compatibility.

⁹ single-sided screen

¹⁰ double-sided screen

¹¹ Complete one section for each processor.

QUALITY ASSURANCE PROGRAM 12

MEDICAL HISTORY

Is a patient history taken as part of the mammographic study? Y N

ı

If yes, indicate which of the following are included in the medical history:

current breast symptoms (mass, nipple discharge, dimpling, etc.)

family history (mother, sister, pre- or post-menopausal)

age at onset and upon cessation of menses

present menstrual status (pre-menopausal, peri-menopausal, post-menopausal)

hormonal pharmaceuticals (birth control pills, menopausal symptoms, fertility drugs, etc., and when)

prior malignancies (type and therapy)

previous mammogram (when and where)

benign breast disease

age at first live birth

nulliparity

any previous breast surgery (type, at what age)

¹² The Q.A. program must meet the requirements of NAC Ch. 457 and 21 CFR 900.

	MAMMOGRAM REPORT						
Does the report include all finding	gs, mammographic/radiographic interpretation a	and clinical/medical examination?					
N/A (no clinical exam done)	No, mammographic findings on	ly Yes, all findings reported					
In the absence of mammographic	In the absence of mammographic findings, is a recommendation made on the basis of the clinical/medical examination?						
N/A (no clinical exam done)	No, mammographic findings on	ly Yes, all findings reported					
What mechanism is in place for received the report? 13	follow-up of positive or equivocal results, and	d to assure that the patient's physician has					
• • • • • • • • • • • • • • • • • • •	, provide a copy of the written authorization for aphy without a prescription signed by a licensed						
Who receives reports for self-refe	rred patients? (check all that apply)						
Each patient must supply the	name of a care physician care provider.						
Patient may select a physiciar	n from an available pool to receive the report. 1	4					
N/A — all patients are referre	d by physicians.						
Which of these follow-up mechan	isms are in place? (check all that apply)						
results of biopsies	cancers with negative mammograms	cancers with positive mammograms					
minimal cancers detected	cancers detected by mammography alone	cancers detected by physical exam alone					
cancers detected by both mar	mmography and physical exam man	nmographic localizations with positive results					
positive mammograms withou	ut evidence of cancer						
In what proportion of cases are a	dditional views done? % Additio	onal imaging exam tests ordered? %					
Specify the types of additional ex	ams ordered:						
What is your facility's current film	retention policy? years In what form	n are results obtained?					
-	· · · · · · · · · · · · · · · · · · ·	FILM, MICROFILM COPIES, ETC.					
What is done with the films and r	eports after this time period?						

¹³ NAC 457.345(4)(b)¹⁴ Provide copy of current list with application.

I attest that the information provided	d in this application is accurate and comp	plete to the best of my know	wledge.
NAME	TITLE	SIGNATURE	DATE
Current fee: \$551.00 per machine.	Fee is non refundable Per NAC 457.295.	Make check payable to: N	levada State Health Divisior

Applications that have SATISFIED ALL REQUIREMENTS may take up to two weeks for processing.

A valid certificate must be posted prior to operation of the mammography machine.

RADIOLOGIST INFORMATION/CHANGES

	currently provides r	adiological services to	
NAME OF RADIOLOGY GROUP			NAME OF FACILITY
supervision of the quality assurar	ice program, the radiological	physicist and the mammogra	nterpretation of mammograms, and aphers involved in this program. This e of the mammographers as well as
If at any time	no longe	r provides these services to	
NAME OF RAD	IOLOGY GROUP	-	NAME OF FACILITY
	will notify Nevada	Division of Health within ten (10	0) work days. At that time, any existing
NAME OF CHIEF TECHNOLOGIST OR FACILITY ADM	NISTRATOR		
	uthorization must be returned		rifies eligibility for a Certificate. The e the Radiology Group discontinues
		adiological services to	
NAME OF PHYSICIST OR PHYSICIST GROUP			NAME OF FACILITY
as an integral part of their mamm	ography services.		
If at any time		r provides these services to _	
NAME OF PHYSICIST	OR PHYSICIST GROUP		NAME OF FACILITY
	will notify Nevada	Division of Health within ten (10) work days. At that time the Chief
NAME OF CHIEF TECHNOLOGIST OR FACILITY ADMIN	ISTRATOR		
	ponsibility. The Facility will	enlist the mandated Physic	ssional coverage until a new Physicist ist support before any repairs are
SUPERVISING RADIOLOGIST	ORIGINAL S	SIGNATURE	DATE
PHYSICIST OR PHYSICIST GROUP REPRESENTAT	TVE ORIGINAL S	SIGNATURE	DATE
FACILITY CORPORATE REPRESENTATIVE	ORIGINAL	SIGNATURE	DATE