	Department of			This inspection checklist is the tool OL		
	Ith & Human Services		R432-950 Mamm	nography Quality	Assurance	licensors use to ensure consistency for every inspection. <i>(Revised 10/2024)</i>
Facility Name:		Facility ID:		Phone Number:		Notes / Sticky Notes
Address				Email Address:		
	1		I			
 (М	Please review the following items ark with a check mark if completed and	during the inspe I make and necessa	ection: ary notes)	Pleas (Mark w	se review the following items d vith a check mark if completed and n	uring the inspection: nake and necessary notes)
	Quality Assurance past 12 months (Physician documents annually that they provide oversight)				List of current employees and former employees past 6 months	
	Current Certification of facility				Policies and Procedures (Reviewed annually)	
	Licensure or certification of personnel				Documentation that mammogram unit is safe	
	Documentation that facility has approval by the American College of Radiology (ACR)				Monthly evaluation of image quality	
	Designated Physician Supervisor				Record accessibility	
	Medical Physicist certified by the American Board of Radiology				Education provided to patients	
	List of Radiologic Technologists				Collecting and Reporting Data process	
Inspection Info	rmation:	-			·	
- All areas that are during the inspectio		r this inspection. Du	ring the inspection, the lice	ensor will ask to have locke	ed areas unlocked. All accessible areas	must be compliant with all applicable rules
- I will email you t	his inspection checklist after the inspec	tion is completed.	l will send you an official	inspection report once th	nis inspection has been approved by	OL management.
	oncompliances are documentation and sure compliance maintenance.	/or records, please	submit them to Licensing	g by the correction requir	ed date listed. A licensor may condu	ct a follow-up inspection to verify

- You may submit f	You may submit feedback on this inspection through your Licensing Portal or at: DLBC.utah.gov													
	Signature Information													
Inspection Type:		Date:		Time Started:		Time Ended:								
	Number of rule noncompliances:		Name of Individual Info	ormed of this Inspection:										
	Licensor(s) Conducting this Inspection:				OL Staff Observing Inspection:									
	The Licensor reviewed compliance.	Please sign/t	ype individual informed ı	name and date of review:										

Utah Depa	irtment of	Inspection C	Checklist		This inspection checklist is the tool OL licensors use to
	& Human Services Background Checks	R432-950 Mammograph	y Quality Assura	nce	ensure consistency for every inspection.
		Licensor Introducto	ory ltems	_	
	Introduction of any unknow	n OL staff to the provider			
	Give a brief explanation of t	he inspection process to the provider			
	ASK: the provider if they wa you conduct the walk- thou them.	nt you to tell staff about rule noncompliances as gh, or wait until the inspection is over to tell			
	Wash hands or use hand sa	nitizer before touching items in the facility.			
		General Not	es		

		R	ULES	CHECK	(LIST		
Rule # R432	Rule Description C = Compliant NC = Not Compliant NA = Not Assessed during this inspection		NC	NA	Compliance Required By Date:	Corrected During Inspection	Notes
R432-35-4. Covered Pro	· · · · · ·	с	NC	NA	Date		Notes
R432-35-4(1)	(1) The covered provider shall enter required information into DACS to initiate a certification for direct patient access of each covered individual before issuance of a provisional license, license renewal, or engagement as a covered individual.						
R432-35-4(2)(a)-(b)	 (2) The covered provider shall ensure the engaged covered individual: (a) signs a criminal background screening authorization form that is available for review by the department; and (b) submits fingerprints within 15 working days of engagement. 						
R432-35-4(3)	(3) The covered provider shall ensure DACS reflects the current status of the covered individual within five working days of the engagement or termination.						
R432-35-4(4)	(4) The covered provider may provisionally engage a covered individual while certification for direct patient access is pending as permitted in Section 26B-2-239.						
R432-35-4(5)	(5) If the department determines an individual is not eligible for direct patient access, based on information obtained through DACS and the sources listed in Section R432-35-8, the department shall send a notice of agency action, as outlined in Rule R432-30, to the covered provider and the individual explaining the action and the individual's right of appeal.						
R432-35-4(6)	(6) The covered provider may not arrange for a covered individual who has been determined not eligible for direct patient access to engage in a position with direct patient access.						

R432-35-4(7)	(7) The department may allow a covered individual to have direct patient access with conditions, during an appeal process, if the covered individual demonstrates to the department, the work arrangement does not pose a threat to the safety and health of patients or residents.			
R432-35-4(8)	(8) The covered provider that provides services in a residential setting shall enter required information into DACS to initiate and obtain certification for direct patient access for each individual 12 years of age and older, who is not a resident, and resides in the residential setting. If the individual is not eligible for direct patient access and continues to reside in the setting, the department may revoke an existing license or deny licensure for healthcare services in the residential setting.			
R432-35-4(9)(a)-(d)	 (9) The covered provider seeking to renew a license as a health care facility shall utilize DACS to run a verification report and verify each covered individual's information is correct, including: (a) employment status; (b) address; (c) email address; and (d) name. 			
R432-35-4(10)	(10) An individual or covered individual seeking licensure as a covered provider shall submit required information to the department to initiate and obtain certification for direct patient access before the issuance of the provisional license. If the individual is not eligible for direct patient access, the department may revoke an existing license or deny licensure as a health care facility.			

			R	ULES	CHECKLIST			
Rule # R432-950	Rule Description C = Compliant NC = Not Compliant NA = Not Assessed during this inspection		NC	NA	Compliance Required By Date:	Corrected During Inspection	RISK: Low Moderate High Extreme	Notes
<u>R380-80-4. Providers' l</u>	Duty to Help Protect Clients.	с	NC	NA	Date			Notes
R380-80-4(1)	(1) The provider shall protect each client from abuse, neglect, exploitation, and mistreatment.							
<u>R380-80-5. Provider Co</u>	de of Conduct.	с	NC	NA	Date			
R380-80-5(4)	(4) Each provider shall protect clients from abuse, neglect, harm, exploitation, mistreatment, fraud, and any action that may compromise the health and safety of clients through acts or omissions and shall instruct and encourage others to do the same.							
<u>R432-950-4. Facility Qu</u>	ality Assurance	с	NC	NA	Date			Notes
R432-950-4(1)	The facility shall conduct a quality assurance program to assure the operation and the services provided are in accordance with R432-950.							
R432-950-4(2)	The facility shall correct identified deficiencies to produce desired results.							
R432-950-4(3)	The facility shall evaluate the corrections required for a systems change to update the quality assurance plan.							
<u>R432-950-5. Complian</u>	e with State and Local Rules	с	NC	NA	Date	CDI		Notes
R432-950-5(1)	A supplier of mammography services shall comply with all applicable Federal, State, and local laws and regulations pertaining to radiological services and mammography services							

R432-950-5(2)(a)-(c)	The facility shall maintain documentation showing that it complies with all applicable state and local laws and rules pertaining to radiological and mammography services. This includes the following: (a) Certification of the facility; (b) Licensure or certification of the personnel; (c) Documentation that the facility has been approved by the American College of Radiology (ACR).						
R432-950-6. Facility Over	rsight	с	NC	NA	Date	CDI	Notes
R432-950-6(1)	The facility is responsible for the overall quality of the mammography conducted.						
R432-950-6(2)(a)-(c)	The facility shall have available, either on staff or through arrangement, sufficient qualified staff to meet patients' needs relating to mammography. Sufficient staff includes the following: (a) A designated physician supervisor who meets the requirements for qualified physicians specified by the Utah Department of Commerce; (b) A medical physicist who is certified by the American Board of Radiology in Radiological Physics or Diagnostic Radiological Physics, or who meets the requirements specified by the Department of Environmental Quality; (c) One or more radiologic technologists who meet the requirements specified by the Utah Department of Commerce pursuant to Section 26-21a-203						
<u>R432-950-7. Physician, P</u>	hysicist and Radiologic Technologist Standards	с	NC	NA	Date	CDI	Notes
R432-950-7(1)	A physician interpreting mammograms or supervising mammography, or both, shall provide documentation to the Department upon request showing he meets minimum qualifications specified by the Utah Department of Commerce and the Mammography Quality Standards Act. A qualified physician shall interpret the results of all mammograms. Diagnostic mammography shall be done under the direct on-site supervision of a qualified physician						

R432-950-7(2)(a)-(e)	A radiologic technologist shall meet the following requirements and the facility shall provide documentation to the Department upon request showing the radiologic technologist: (a) Meets minimum qualifications specified by the Utah Department of Commerce and the Mammography Quality Standards Act; (b) Obtains on-the-job training in mammography under the supervision of a qualified physician, or the supervising radiologic technologist, or both; (c) Is competent in breast positioning and compression as determined from critiques by a qualified physician of mammogram films taken by the radiologic technologist; (d) Is knowledgeable in facility policies concerning technical factors, radiation safety, radiation protection, and quality control as evaluated by the radiologic technologist's supervisor; (e) Receives continuous supervision and feedback on image quality from the interpreting or supervising physician.	0					
R432-950-7(3)(a)-(c)	A medical physicist must: (a) be certified in an acceptable specialty by one of the bodies approved by the FDA to certify medical physicists; (b) be licensed or approved by a State to conduct evaluations of mammography equipment as required by State law; or (c) for those medical physicists associated with facilities that apply for accreditation before October 27, 1997, who meet training and experience requirements of Mammography Quality Standards Act and its implementing regulations						
R432-950-8. Personnel	Requirements	С	NC	NA	Date	CDI	Notes
R432-950-8(1)	The facility shall document that new staff orientation and ongoing in-service training is based on current written facility policies and procedures.						
R432-950-8(2)	Personnel shall have access to the facility's written policies and procedures when on duty						

R432-950-8(3)	The facility shall implement a standardized orientation program for each employment position including the time for completing training.						
R432-950-8(4)	A written in-service training program shall identify the topics and frequency of training including an annual review of facility policies and procedures.						
R432-950-8(5)	The facility shall maintain personnel records documenting that each employee is qualified and competent to perform respective duties and responsibilities by means of appropriate licensure or certification, experience, orientation, ongoing in-service training, and continuing education.						
R432-950-8(6)	The facility shall retain personnel records for terminated employees for a minimum of four years following the final date of termination.						
<u>R432-950-9. Equipment :</u>	Standards	с	NC	NA	Date	CDI	Notes
R432-950-9(1)	Mammogram units shall be designed specifically for mammography and shall have a compression device and						
	the capability for placement of a grid.						
R432-950-9(2)	The facility shall maintain current written policies and procedures for operating equipment.						
R432-950-9(2) R432-950-9(3)	The facility shall maintain current written policies and						
	The facility shall maintain current written policies and procedures for operating equipment. Prior to initiating operation of a mammogram unit it shall be registered with the Utah Department of Environmental Quality.				Date		Notes

10(2)	The facility shall maintain documentation that employees have been trained on safety standards for radiation.						
10(3)	The facility shall maintain procedure manuals and logs for equipment quality control.						
10(4)(a-b)	The facility shall maintain documentation that the quality control program complies with ACR quality control manuals for mammography or the equivalent. (a) Equivalent programs shall include a quality control program for equipment, mammogram unit performance, and film processors, approved by the Utah Department of Environmental Quality. (b) Equivalent programs shall contain stated objectives achieved by procedures comparable to objectives and procedures in the American College of Radiology Quality Control Manuals for Mammography.						
10(5)	Accreditation by the American College of Radiology Mammography Program documents compliance with mammogram unit quality control requirements in R432-950-10(1).						
<u>R432-950-11. Technical Spectral Spectrad Spectrad Spectrad Spectrad Spectrad Spectr</u>	pecifications for Mammography	-					
	<u>h </u>	С	NC	NA	Date	CDI	Notes
11(1)	The facility shall have available a phantom for use in the facility's ongoing quality control program.				Date	СЫ	Notes
11(1)	The facility shall have available a phantom for use in the				Date		Notes

<u>R432-950-12. Physicia</u>	n Supervisor Responsibility	с	NC	NA	Date	CDI	Notes
12(1)(a-e)	A physician supervisor is responsible for general oversight of the quality control program of the facility. Oversight responsibilities include: (a) Annual review of the policy and procedure manual; (b) Verification that the equipment and facility personnel meet applicable federal, state and local licensure and registration requirements; (c) Verification that equipment is performing properly; (d) Verification that safe operating procedures are used to protect facility personnel and patients; (e) Verification that all other requirements of R432-950 are being met.						
12(2)	The physician shall document annually that he provides oversight for the quality control of the mammography service.						
<u>R432-950-13. Mammo</u>	graphy Records	с	NC	NA	Date	CDI	Notes

13(1)(a-e)	 A medical record shall be maintained for each patient on whom screening or diagnostic mammography is performed. (a) Provision shall be made for the filing, safe storage and accessibility of medical records. (b) Records shall be protected against loss, defacement, tampering, fires, and floods. (c) Records shall be protected against access by unauthorized individuals. (d) All records shall be readily available upon the request of: (i) The attending physician, (ii) Authorized representatives of the Department for determining compliance with licensure rules; (iii) Any other person authorized by written consent. (e) The facility shall establish a system to assure that the patient's mammogram is accessible for clinical follow-up when requested. (i) A copy of the mammogram and other appropriate information shall be sent to the requesting party responsible for subsequent medical care of the patient no later than 14 working days from the request for information. This shall include the full notification and follow up required under Utah Code 26-21a-206 and Administrative Code R432-950-14. (ii) Medical information may be released only upon the written consent of the patient of her legal representative. 				
13(2)	The facility shall attempt to obtain a prior mammogram for each patient if the prior mammogram is necessary for the physician to properly interpret the current exam.				
13(3)(a-d)	 The interpreting physician shall prepare and sign a written report of his interpretation of the results of the screening mammogram. (a) The written report shall include a description of detected abnormalities and recommendations for subsequent follow-up studies. (b) The interpreting physician shall render the report as soon as reasonably possible. (c) The interpreting physician or his designee shall document and communicate the results of the report to the referring physician or his designated representative by telephone, by certified mail, or in such a manner that receipt of the report is assured. (d) The interpreting physician or his designee shall notify self-referred patients, that is, patients who have no referring physician, of the results of the screening study in writing and in lay language. 	0			

13(4)	The interpreting physician or his designee shall document and communicate the results of all diagnostic reports in the high probability category with suspicion of breast cancer to the referring physician or his designated representative by telephone, by certified mail, or in such a manner that receipt of the report is assured.						
13(5)(a-b)	The physician shall document and communicate in person in lay language, by certified mail, or in such a manner that receipt of the diagnostic report is assured to all self-referred patients within the high probability category with a suspicion of breast cancer. The report shall indicate whether the patient needs to consult with a physician. (a) The interpreting physician or his designee shall attempt to make a follow-up contact with the patient to determine whether she has consulted a physician for follow-up care. (b) The interpreting physician or his designee shall document in the patient's medical record attempts to communicate the results to the patient.						
13(6)	The facility shall retain the original and subsequent mammograms for a period of at least five years from the date of the procedure.						
<u>R432-950-14. Educatio</u>	n and Notification Requirements	с	NC	NA	Date	CDI	Notes
14(1)	A patient has the right to be treated with dignity and afforded privacy during the examination.						
14(2)(a-c)	The facility shall establish an education system to ensure that the patient understands: (a) The purpose of the mammogram and how it is used to screen for breast cancer; (b) The process required to obtain the mammogram; (c) The importance of the screening mammography to her ongoing health.						

14(3)	As required in Utah Code 26-21a-206, the facility shall include the following notification and information with a mammography result provided to a patient with dense breast tissue: "Your mammogram indicates that you have dense breast tissue. Dense breast tissue is common and is found in as many as half of all women. However, dense breast tissue can make it more difficult to fully and accurately evaluate your mammogram and detect early signs of possible cancer in the breast. This information is being provided to inform and encourage you to discuss your dense breast tissue and other breast cancer risk factors with your health care provider. Together, you can decide what may be best for you. A copy of your mammography report has been sent to your health care provider. Please contact them if you have any questions or concerns about this notice."						
14(4)	The copy of the mammography report provided to the patient and the health care provider shall include the dense breast tissue notification required under Utah Code 26-21a-206.						
<u>R432-950-15. Collecting</u>	R432-950-15. Collecting and Reporting Data		NC	NA	Date	CDI	Notes
15(1)(a-c)	 The facility shall establish a system for collecting and periodic reporting of mammography examinations and clinical follow-up as provided below: (a) Clinical follow-up data shall include the follow-up on the disposition of positive mammographic findings, and the correlation of the surgical biopsy results with mammogram reports. (b) The facility shall maintain records correlating the positive mammographic findings to biopsies done and the number of cancers detected. (c) The facility shall report the results of the outcomes annually to the Department or its designated agent, on forms furnished by the Department. The report shall include as a minimum: (i) The number of patients receiving screening mammograms; (ii) Total number of patients diagnosed with breast cancer based on a screening mammogram; (iv) The number and names of individuals with positive mammographic findings to biopsed on patients receiven the partment on the partment of the partment for biopsy based on a screening mammogram; (iv) The number of patients diagnosed with breast cancer based on a screening mammogram; (iv) The number and names of individuals with positive mammographic findings lost to follow-up. 	0	0				
15(2)	The Department or its designated agent shall provide each reporting facility, on a schedule determined by the Department, summary statistical reports which permit each facility to compare its results to statewide and other comparative statistics.						

<u>R432-950-16. State Ce</u>	R432-950-16. State Certification		NC	NA	Date	CDI	Notes
16(1)	No facility, person or governmental unit acting severally or jointly with any other person, may establish, conduct or maintain a mammography unit without first obtaining a state certificate from the Department.						
16(2)	An applicant for state certification shall file a Request for Agency Action/Certification Application with the Utah Department of Health on forms furnished by the Department.						
16(3)(a-c)	Each facility shall comply with all zoning, building and licensing laws, rules and ordinances and codes of the city and county in which the facility is located. The applicant shall submit the following to the Department: (a) Verification of participation and quality control by the American College of Radiology for monitoring mammography services in the facility; (b) Verification of licensure or certification of required personnel; (c) Fees established by the Utah State Legislature pursuant to Section 63-38-3.						
16(4)(a-b)	The Department shall render a decision on the initial certification within 60 days of receipt of a completed application packet or within 6 months of date that the first component of an application packet was received. (a) Upon verification of compliance with state certification requirements, the Department shall issue a provisional certificate. (b) The Department shall issue a notice of agency decision under the procedures for informal adjudicative proceedings denying a state certification if the applicant is not in compliance with the applicable laws or rules. The notice shall state the reasons for denial.						

16(5)(b-d) should be [a-c]	Certificate Contents and Provisions. The state certificate shall include the name of the mammography facility, owner, supervising physician, address, issue and expiration dates of the state certificate and the certificate number. (b) The state certificate may be issued only to the owner and for the premises described in the application and shall not be assignable or transferable. (c) Each state certificate is the property of the Department and shall be returned within five days if the certification is suspended, revoked, or if the operation of the facility is discontinued. (d) The state certificate shall be prominently displayed where it can be easily viewed by the public.				
16(6)(a-b)	Certification periods shall be for 24 months, and expire at midnight 24 months from the date of issuance. (a) A request for renewal and applicable fees shall be filed with the Department 15 days before the state certificate expires. (b) Failure to make a timely renewal shall result in assessment of late fees as established by the Utah State Legislature pursuant to Section 26-21a-203.				
16(7)(a-c)	The owner shall submit a Request for Agency Action/Application to amend or modify state certification status at least 30-days before any of the following proposed or anticipated changes occur: (a) Change in the name of the facility; (b) Change in the supervising physician; (c) Change in the owner of the facility.				
16(8)(a-c)	The owner who wants to cease operation shall complete the following: (a) Notify the patients within 30 days before the effective date of closure. (b) Make adequate provision for the safekeeping of records and notify the department where those records will be stored. (c) Return the state certificate to the Department within five days after the facility ceases operation.				

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16(9)(a-b)	 The Department may issue a provisional state certificate to a facility as an initial certification and may issue a provisional state certification to a facility that does not fully comply with the requirements for a standard certification but has made acceptable progress towards meeting the requirements. (a) In granting a provisional state certification, the Department must be assured that the lack of full compliance does not harm the health, safety, and welfare of the patients. (b) A provisional state certificate is nonrenewable and shall be issued for no more than 6 months. 							
<u>R432-950-17. Inspecti</u>	n <u>s</u>	с	NC	NA	Date	CDI		Notes
Upon presentation of proper identification, authorized representatives of the Departme including medical records, when it is determined by the Department to be necessary to			iall be rtain c	allow ompli	ed to enter a fac ance with state l	tility at any rea law and rules p	sonable time promulgated	without a warrant and be permitted to review records under Section 26-21a-205.
17(1)	Each facility may be inspected by the Department or its designee to determine compliance with minimum standards and the applicable rules.							
17(2)	Upon receipt of the survey results of the ACR, the facility shall submit copies of the certificate and the survey report and recommendations.							
17(3)	The accreditation documents are open to the public.							
17(4)	The Department may conduct periodic validation inspections of facilities accredited by the ACR for the purpose of determining compliance with state requirements.							
R432-950-18. Enforcement and Appeal Process			NC	NA	Date	CDI		Notes
R432-950-18	Whenever the Department has reason to believe that the facility is in violation of Section 26-21a-203 or any of the rules adopted pursuant to Title 26, Chapter 21, the Department shall issue a written Statement of Findings/Plan of Correction to the certified facility.							