

# Relocation Assessment Evidence Submission for Distance Review

## DIAGNOSTIC IMAGING

### Introduction

The facility's evidence submission will be utilized to determine if the facility meets the DAP relocation assessment accreditation standards. The facility's current accreditation award will be transferred to the new address/location after all requirements have been implemented.

The following sections outline the criteria that will be assessed by evidence submission. Follow-up may be required during the relocation assessment visit to determine if the evidence is implemented.

FACILITY INFORMATION	
Facility name:	_____
New address:	_____
City: _____	Postal code: _____
Imaging service phone number: _____	Projected date of new facility opening: _____

CONTACT PERSON FOR IMAGING SERVICE ACCREDITATION ACTIVITIES	
Name: _____	Title: _____
Address: _____	
City: _____	Postal code: _____
Phone: _____	Fax: _____
Cellular phone: _____	Email: _____

Relocation Assessment Evidence Submission for Distance Review *continued*

Facility name: \_\_\_\_\_

<b>MEDICAL STAFF</b>							
Criteria and descriptors Note: M indicates mandatory.			Guidance for evidence submission	Evidence attached	Not applicable	No evidence	Explain
<b>DMS1.2</b>	<b>Medical leaders must attend the diagnostic service to assess the quality and safety of service.</b>		Log of the medical leaders visits to the facility/service, if remotely supervised.				
	DMS1.2.1- DMS1.2.15	<b>M</b> The medical leader visits the facility prior to assuming responsibility for medical leadership of a new service.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	

<b>HUMAN RESOURCES</b>							
Criteria and descriptors Note: M indicates mandatory.			Guidance for evidence submission	Evidence attached	Not applicable	No evidence	Explain
<b>DHR5.2</b>	<b>Orientation and ongoing training is provided to existing staff to uphold the quality and safety of the diagnostic service.</b>		Orientation records to the new space/facility and policies or procedures that have changed as a result of the relocation (e.g. emergency evacuation procedures, codes, etc.).  Orientation records on any new equipment (both imaging systems and ancillary equipment).				
	DHR5.2.1	<b>M</b> Orientation and training is provided to existing staff in response to changing roles, technology, competency demands, laws and regulations or after an extended leave.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	DHR5.2.3	<b>M</b> Existing staff are provided with ongoing training or orientation in imaging system and ancillary equipment use, maintenance and safety features.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	

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Facility name: \_\_\_\_\_

GENERAL SAFETY								
Criteria and descriptors Note: M indicates mandatory.			Guidance for evidence submission	Evidence attached	Not applicable	No evidence	Explain	
DSA1.2	<b>A safety manual is readily available to staff that includes:</b>		Safety manual, highlighting the procedures updated following the relocation (e.g. evacuation plans, emergency response procedures, staff working alone or in isolation).					
	DSA1.2.1	M		<ul style="list-style-type: none"> <li>how to access first aid services and/or medical assistance for staff related injuries</li> </ul>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	DSA1.2.4	M		<ul style="list-style-type: none"> <li>requirements for use of personal protective and other safety equipment</li> </ul>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	DSA1.2.5	M		<ul style="list-style-type: none"> <li>Workplace Hazardous Materials Information System (WHMIS) program information</li> </ul>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	DSA1.2.6	M		<ul style="list-style-type: none"> <li>emergency evacuation plans</li> </ul>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	DSA1.2.7	M		<ul style="list-style-type: none"> <li>procedures to protect staff "working alone" or in "isolation"</li> </ul>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	DSA1.2.8	M		<ul style="list-style-type: none"> <li>procedures to manage violent and aggressive behaviour</li> </ul>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
DSA2.1	<b>The design and layout of the physical space meets laws, regulations and codes.</b>							
	DSA2.1.1	M	A professional engineer, responsible for the build, has attested that the new construction or structural changes meet the minimum CSA standards.	City-issued occupancy permit.	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	

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Facility name: \_\_\_\_\_

<b>PATIENT SAFETY</b>							
Criteria and descriptors Note: M indicates mandatory.			Guidance for evidence submission	Evidence attached	Not applicable	No evidence	Explain
DPS6.1	There are procedures to handle medical emergencies in a timely and effective manner.		Updated emergency medical response protocol.				
	DPS6.1.1	M There is a medical emergency response protocol in place.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	

<b>INFECTION PREVENTION AND CONTROL</b>							
Criteria and descriptors Note: M indicates mandatory.			Guidance for evidence submission	Evidence attached	Not applicable	No evidence	Explain
DIPC5.1	There is a defined follow up process that addresses possible or actual blood and body fluid exposures.		Updated blood and body fluid exposure procedures.				
	DIPC5.1.3	M There are documented policies and procedures for the prevention and follow-up of blood and body fluid exposures.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	

Relocation Assessment Evidence Submission for Distance Review *continued*

Facility name: \_\_\_\_\_

<b>EQUIPMENT AND SUPPLIES</b>							
<b>Criteria and descriptors</b> Note: M indicates mandatory.			<b>Guidance for evidence submission</b>	<b>Evidence attached</b>	<b>Not applicable</b>	<b>No evidence</b>	<b>Explain</b>
<b>DES2.1</b>	<b>Acceptance testing is performed after purchase and prior to clinical use of equipment.</b>						
	DES2.1.4	<b>M</b> Acceptance testing of imaging equipment includes an initial inspection of the equipment and any ancillary equipment.	Acceptance reports for any new ancillary equipment (e.g. pumps, lead aprons & lead shields).	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>DES3.5</b>	<b>Quality control procedures are established and used to monitor performance of electronic display devices (monitors/image display systems).</b>		Calibration/ acceptance testing reports for any new primary or secondary displays (monitors).				
	DES3.5.1	<b>M</b> The performance of all new electronic display devices used for the interpretation of diagnostic images and guidance during interventional procedures is tested to verify performance prior to clinical use.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	

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Facility name: \_\_\_\_\_

<b>RADIATION SAFETY</b>							
<b>Criteria and descriptors</b> Note: M indicates mandatory.			<b>Guidance for evidence submission</b>	<b>Evidence attached</b>	<b>Not applicable</b>	<b>No evidence</b>	<b>Explain</b>
<b>RS4.2</b>	<b>New and replaced medical X-ray equipment is registered with the Diagnostic Accreditation Program of BC.</b>		DAP equipment registration forms for all new or replaced equipment.  <i>Note: Forms can be found at:</i> <a href="https://www.cpsbc.ca/programs/dap/accreditation/diagnostic-imaging">https://www.cpsbc.ca/programs/dap/accreditation/diagnostic-imaging</a>				
	RS4.2.1	<b>M</b> New and replaced medical X-ray equipment is registered with the Diagnostic Accreditation Program of BC and includes the facility name and address.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>RS5.1</b>	<b>Radiation protection surveys are conducted to assess safety when:</b>		Radiation protection survey report for all X-ray, CT and Mammography rooms.				
	RS5.1.1	<b>M</b> • there is a new installation		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>RS6.1</b>	<b>Appropriate steps are taken to ensure adequate shielding is present in controlled and uncontrolled areas.</b>		Shielding plans.				
	RS6.1.1	<b>M</b> The radiation levels in controlled areas that are occupied routinely by radiation workers are such that no radiation worker is occupationally exposed to more than 20 mSv per year.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	

Relocation Assessment Evidence Submission for Distance Review *continued*

Facility name: \_\_\_\_\_

<b>RADIOLOGY</b>							
<b>Criteria and descriptors</b> Note: M indicates mandatory.			<b>Guidance for evidence submission</b>	<b>Evidence attached</b>	<b>Not applicable</b>	<b>No evidence</b>	<b>Explain</b>
<b>RA3.3</b>	<b>Examinations are performed following established protocols.</b>		If new imaging equipment is installed, protocols updated to reflect the system(s) in place.				
	RA3.3.1	<b>M</b> Protocols are readily available to staff performing the examination.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	RA3.3.3	<b>M</b> There are protocols for the pediatric population.	Technique charts are reflective of the imaging system(s) in place.	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	RA3.3.8	<b>M</b> Technique charts are available and reflective of the equipment used.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>RA12.1</b>	<b>Acceptance testing is performed after purchase and prior to clinical use of film-based systems.</b>		Acceptance testing report for film-based systems.				
	Acceptance testing includes visual and functional testing of the:						
	RA12.1.1	<b>M</b> • mechanical properties		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	RA12.1.2	<b>M</b> • safety systems		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>RA12.2</b>	<b>Acceptance testing is performed after purchase and prior to clinical use of CR/DR systems that includes:</b>		Acceptance testing report for CR/DR radiographic systems (and mobile units).				
	Acceptance testing includes visual and functional testing of the:						
	RA12.2.1	<b>M</b> • mechanical properties	<i>Note: Acceptance testing is not required for mobile units relocated within the same facility.</i>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	RA12.2.2	<b>M</b> • safety systems		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	

# Relocation Assessment Evidence Submission for Distance Review *continued*

Facility name: \_\_\_\_\_

RADIOLOGY							
Criteria and descriptors Note: M indicates mandatory.			Guidance for evidence submission	Evidence attached	Not applicable	No evidence	Explain
RA12.3	<b>Acceptance testing is performed after purchase and prior to clinical use of radioscopic systems that includes:</b>		Acceptance testing report for radiographic systems (including film-based and mobile units).				
	Acceptance testing includes visual and functional testing of the:						
	RA12.3.1	<b>M</b>	• mechanical properties	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	RA12.3.2	<b>M</b>	• safety systems	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
			<i>Note: Acceptance testing is not required for mobile units relocated within the same facility.</i>				



Relocation Assessment Evidence Submission for Distance Review *continued*

Facility name: \_\_\_\_\_

<b>MAMMOGRAPHY</b>							
<b>Criteria and descriptors</b> Note: M indicates mandatory.			<b>Guidance for evidence submission</b>	<b>Evidence attached</b>	<b>Not applicable</b>	<b>No evidence</b>	<b>Explain</b>
<b>MA3.3</b>	<b>Examinations are performed following established protocols.</b>		If new imaging equipment is installed, protocols updated to reflect the system(s) in place.				
	MA3.3.1	<b>M</b> Protocols are readily available to staff performing the examination.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>MA12.1</b>	<b>Acceptance testing of film-based systems is performed by a medical physicist after purchase and prior to clinical use.</b>		Physicist's acceptance testing report for film-based mammography systems.				
	Acceptance testing includes visual and functional testing of the:						
	MA12.1.1	<b>M</b> • mechanical properties		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	MA12.1.2	<b>M</b> • safety systems		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>MA12.2</b>	<b>Acceptance testing of CR/DR systems is performed by a medical physicist after purchase and prior to clinical use of mammography X-ray equipment.</b>		Physicist's acceptance testing report for CR/DR mammography systems.				
	Acceptance testing includes visual and functional testing of the:						
	MA12.2.1	<b>M</b> • mechanical properties		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	MA12.2.2	<b>M</b> • safety systems		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	

Relocation Assessment Evidence Submission for Distance Review *continued*

Facility name: \_\_\_\_\_

ULTRASOUND							
Criteria and descriptors Note: M indicates mandatory.			Guidance for evidence submission	Evidence attached	Not applicable	No evidence	Explain
US3.3	<b>Examinations are performed following established protocols.</b>		If new imaging equipment is installed, protocols updated to reflect the system(s) in place.				
	US3.3.1	<b>M</b> Protocols are readily available to staff performing the examination.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
US12.1	<b>Acceptance testing is performed after purchase and prior to clinical use of equipment that includes:</b>		Acceptance testing report for each ultrasound system.				
	US12.1.1	<b>M</b> <ul style="list-style-type: none"> <li>visual inspection of system and probes</li> </ul>		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	US12.1.2	<b>M</b> <ul style="list-style-type: none"> <li>electrical leakage current testing of probes</li> </ul>	<i>Note: Ultrasound systems relocated within the same facility do not require acceptance testing.</i>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	

Relocation Assessment Evidence Submission for Distance Review *continued*

Facility name: \_\_\_\_\_

ECHOCARDIOGRAPHY							
Criteria and descriptors Note: M indicates mandatory.			Guidance for evidence submission	Evidence attached	Not applicable	No evidence	Explain
EC3.3	<b>Examinations are performed following established protocols.</b>		If new imaging equipment is installed, protocols updated to reflect the system(s) in place.				
	EC3.3.1	<b>M</b> Protocols are readily available to staff performing the examination.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
EC12.1	<b>Acceptance testing is performed after purchase and prior to clinical use of equipment that includes:</b>		Acceptance testing report for each echocardiography system.				
	US12.1.1	<b>M</b> <ul style="list-style-type: none"> <li>visual inspection of system and probes</li> </ul>		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	US12.1.2	<b>M</b> <ul style="list-style-type: none"> <li>electrical leakage current testing of probes</li> </ul>	<i>Note: Echocardiography systems relocated within the same facility do not require acceptance testing.</i>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	

# Relocation Assessment Evidence Submission for Distance Review *continued*

Facility name: \_\_\_\_\_

<b>COMPUTED TOMOGRAPHY</b>							
<b>Criteria and descriptors</b> Note: M indicates mandatory.			<b>Guidance for evidence submission</b>	<b>Evidence attached</b>	<b>Not applicable</b>	<b>No evidence</b>	<b>Explain</b>
<b>CT3.3</b>	<b>Examinations are performed following established protocols.</b>		If new imaging equipment is installed, protocols updated to reflect the system(s) in place.				
	CT3.3.2	<b>M</b> Protocols are equipment specific.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>CT12.1</b>	<b>Acceptance testing is performed after purchase and prior to clinical use of the equipment.</b>		Acceptance testing report for each CT scanner.				
	Acceptance testing includes visual and functional testing of the:						
	CT12.1.1	<b>M</b> • mechanical properties		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	CT12.1.2	<b>M</b> • safety systems		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	

# Relocation Assessment Evidence Submission for Distance Review *continued*

Facility name: \_\_\_\_\_

<b>MAGNETIC RESONANCE IMAGING</b>							
<b>Criteria and descriptors</b> Note: M indicates mandatory.			<b>Guidance for evidence submission</b>	<b>Evidence attached</b>	<b>Not applicable</b>	<b>No evidence</b>	<b>Explain</b>
<b>MR3.3</b>	<b>Examinations are performed following established protocols.</b>		If new imaging equipment is installed, protocols updated to reflect the system(s) in place.				
	MR3.3.3	<b>M</b> Protocols are equipment specific.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>MR12.1</b>	<b>Acceptance testing is performed by a medical physicist after purchase and prior to clinical use of the MRI system.</b>		Acceptance testing report for each MRI system.				
	Acceptance testing procedures include, but are not limited to the following:						
	MR12.1.1	<b>M</b> <ul style="list-style-type: none"> <li>assessment and identification of the fringe fields</li> </ul>		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	MR12.1.2	<b>M</b> <ul style="list-style-type: none"> <li>magnetic field homogeneity</li> </ul>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>		

# Relocation Assessment Evidence Submission for Distance Review *continued*

Facility name: \_\_\_\_\_

<b>MAGNETIC SAFETY</b>							
<b>Criteria and descriptors</b> Note: M indicates mandatory.			<b>Guidance for evidence submission</b>	<b>Evidence attached</b>	<b>Not applicable</b>	<b>No evidence</b>	<b>Explain</b>
<b>MRS1.1</b>	<b>Individuals knowledgeable in MRI safety are involved in planning and review of facility design plans for a new MRI installation.</b>		Facility floor plan showing all four zones.				
	MRS1.1.1	<b>M</b> Any new facility has incorporated the ACR 4 Zone Configuration into their design plans.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>MRS4.3</b>	<b>Education is provided to:</b>		MR Safety training records for non-MRI personnel that may enter Zone 3 or 4.				
	MRS4.3.1	<b>M</b> <ul style="list-style-type: none"> <li>Housekeeping staff</li> </ul>		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	MRS4.3.2	<b>M</b> <ul style="list-style-type: none"> <li>Municipal emergency response staff</li> </ul>		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	MRS4.3.3	<b>M</b> <ul style="list-style-type: none"> <li>Security staff</li> </ul>		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	

Relocation Assessment Evidence Submission for Distance Review *continued*

Facility name: \_\_\_\_\_

<b>NUCLEAR MEDICINE</b>							
<b>Criteria and descriptors</b> Note: M indicates mandatory.			<b>Guidance for evidence submission</b>	<b>Evidence attached</b>	<b>Not applicable</b>	<b>No evidence</b>	<b>Explain</b>
<b>NM3.3</b>	<b>Examinations are performed following established protocols.</b>		If new imaging equipment is installed, protocols updated to reflect the system(s) in place.				
	NM3.3.2	M Protocols are equipment specific.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>NM12.1</b>	<b>Acceptance testing is performed by a medical physicist after purchase and prior to clinical use of gamma camera systems.</b>		Acceptance testing report of each nuclear medicine gamma camera.				
	NM12.1.1	M • multiple-window registration		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	NM12.1.2	M • maximum count rate		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>NM12.2</b>	<b>Acceptance testing is performed after purchase and prior to clinical use of well counter systems.</b>		Acceptance testing report of well counter system.				
	NM12.2.1	M • crystal energy resolution		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	NM12.2.2	M • linear geometry and sensitivity		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	NM12.2.3	M • minimum/maximum detectable levels		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>NM12.3</b>	<b>Acceptance testing is performed after purchase and prior to clinical use of uptake probe systems.</b>		Acceptance testing report for uptake probe.				
	NM12.3.1	M • crystal energy resolution		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>NM12.4</b>	<b>Acceptance testing is performed after purchase and prior to clinical use of dose calibrator systems.</b>		Acceptance testing report for dose calibrator(s).				
	NM12.4.1	M • geometrical sensitivity		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	

# Relocation Assessment Evidence Submission for Distance Review *continued*

Facility name: \_\_\_\_\_

<b>NUCLEAR MEDICINE</b>							
<b>Criteria and descriptors</b> Note: M indicates mandatory.			<b>Guidance for evidence submission</b>	<b>Evidence attached</b>	<b>Not applicable</b>	<b>No evidence</b>	<b>Explain</b>
<b>NM12.5</b>	<b>Acceptance testing is performed after purchase and prior to clinical use of SPECT/CT hybrid systems.</b>		Radiation scatter measurement report.				
	NM12.5.1	<b>M</b> For all SPECT/CT hybrid systems, the radiation levels are monitored at critical areas in the imaging room (e.g. bedside, doorway, workstation, etc.).		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	



Relocation Assessment Evidence Submission for Distance Review *continued*

Facility name: \_\_\_\_\_

<b>BONE DENSITOMETRY</b>							
<b>Criteria and descriptors</b> Note: M indicates mandatory.			<b>Guidance for evidence submission</b>	<b>Evidence attached</b>	<b>Not applicable</b>	<b>No evidence</b>	<b>Explain</b>
<b>BD3.3</b>	<b>Examinations are performed following established protocols.</b>		If new imaging equipment is installed, protocols updated to reflect the system(s) in place.				
	BD3.3.3	<b>M</b> Protocols are readily available to staff performing the examination.		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>BD 11.2</b>	<b>The cross-calibration of the bone densitometry equipment is performed according to standard protocols and prior to clinical use.</b>		Cross-calibration report if the existing system is replaced with: - new hardware - new BMD system				
	Acceptance testing includes visual and functional testing of the:						
	BD12.1.1	<b>M</b> • mechanical properties		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	BD12.1.2	<b>M</b> • safety systems		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>BD 12.1</b>	<b>Acceptance testing is performed after purchase and prior to clinical use of equipment.</b>		Acceptance testing report for bone densitometry system(s).				
	Acceptance testing includes visual and functional testing of the:						
	BD12.1.1	<b>M</b> • mechanical properties		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
	BD12.1.2	<b>M</b> • safety systems		<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	