



Accreditation Program Guide

Diagnostic Imaging:
New Facility
New Imaging Service (modality)
Relocation

Version: September 2018 – v2

TABLE OF CONTENTS

1.0 PURPOSE OF ACCREDITATION 3

2.0 COLLEGE OF PHYSICIANS AND SURGEONS OF ALBERTA (CPSA) ACCREDITATION PROGRAM..... 3

2.1 CPSA Lines of Business 3

2.2 Mission, Vision and Values 4

2.3 CPSA Organizational Structure - Figure 1 5

2.4 Accreditation Program History 6

2.5 Authority and Oversight 6

2.6 Overview of Diagnostic Imaging (DI) Accreditation Program 7

2.7 Personnel..... 8

 2.7.1 CPSA Diagnostic Imaging Accreditation Personnel and Roles..... 8

 2.7.2 Advisory Committee on Diagnostic Imaging (ACDI)..... 8

2.8 Assessment Teams 9

 2.8.1 Assessment Coordinator 9

 2.8.2 Assessor (Quality Assessors and Modality Specific)..... 9

 2.8.3 Physician Reviewer 9

 2.8.3 Team Selection..... 10

 2.8.4 Assessment Team Training 10

 2.8.5 Conflict of Interest / Confidentiality Agreements /Liability..... 10

3.0 STANDARDS DOCUMENT 11

3.1 Standards Overview 11

3.2 Format of Standards..... 12

3.3 Assessment of Compliance (AOC) 13

3.4 Terms and Definitions 14

3.5 Reference Listing 14

3.6 Review and Revision of Standards..... 15

4.0 ACCREDITATION PROCESS – NEW/RELOCATED DI FACILITY ACCREDITATION 16

4.1 Initiation – New/Relocated DI facility..... 17

4.2 Pre-assessment..... 17

4.3 On-site assessment..... 21

4.4 Post-assessment 25

5.0 HONORARIA AND EXPENSE REIMBURSEMENT 31

6.0 ASSESSMENT FEES 31

7.0 LIST OF ACCREDITED DIAGNOSTIC IMAGING DI FACILITIES..... 31

8.0 EXAMINATION/IMAGE REVIEW 32

8.1 Public Imaging Facilities..... 33

8.2 Private Imaging Facilities 33

8.3 Minimum Number of Examinations 34

1.0 Purpose of Accreditation

Accreditation is defined as the public recognition of quality achievement by a healthcare organization, as demonstrated through an independent external peer comparison of the organization's performance against current best practices.

The College of Physicians and Surgeons of Alberta (CPSA) diagnostic accreditation programs:

- assist facilities with a process of ensuring accuracy and reliability of examination/services
- provide standards of practice and assess compliance to these standards
- identify deficiencies that affect the quality of examination/services, and impact patient and/or staff safety
- evaluate a facility's quality system's ability to identify and mitigate risk and variability in system processes
- gives formal recognition of a facility's provision of quality diagnostic services
- encourage and facilitate peer review
- provide educational opportunities for both the facility being accredited and the assessment team
- promote uniformity in practice provincially, where variations in practice are counter-productive for the province
- maintain a comprehensive data repository for scope of service/levels of imaging and resources
- promote standardization and educational initiatives across Canada through inter-provincial collaboration
- promote and encourage dialogue amongst stakeholders on best-practices and best ways to incorporate them into the workflow
- ensure effective medical direction over medical practices so that business interests do not determine the standards of care

2.0 College of Physicians and Surgeons of Alberta (CPSA) Accreditation Program

2.1 CPSA Lines of Business

The CPSA is mandated by legislation to regulate the practice of medicine in Alberta and is responsible for licensing physicians, administering standards of practice and conduct, and resolving physician-related complaints.

It also provides leadership and direction on issues of importance to the health care system such as access to services, quality improvement, patient safety and privacy.

The Council of the CPSA is composed of physicians elected by members of the profession in Alberta, the two Deans of Medicine in Alberta and four members of the public appointed by the Minister of Health and Wellness.

The lines of business for the CPSA are as follows:

- Register physicians
- Investigate and resolve physician-related complaints
- Provide clinical review
- Accredite health facilities
- Guide professional conduct and ethical behavior
- Contribute to public policy affecting health care delivery

2.2 Mission, Vision and Values

Our Mission

Serving the public by guiding the medical profession

Our Vision

Albertans are healthier because the College of Physicians & Surgeons of Alberta:

- ensures that physicians are competent throughout their careers;
- supports physicians in providing compassionate, caring and ethical services to the people of Alberta;
- fosters quality health care for all Albertans through innovation, collaboration and cooperation with other key stakeholders; and
- advocates for public policy that contributes to the health of Albertans.

Our Values

We do the right thing.

We act responsibly, respectfully and with integrity, aspiring to be fair and responsible. We acknowledge our mistakes as well as our successes, and strive to do what's right in the service to the public.

We make informed decisions.

Our decisions are based on evidence, knowledge, experience and best practice. We plan, measure outcomes and apply what we learn.

We empower people.

We believe people perform best when they see the Vision, set their own goals, have the resources they need and aspire to excellence and personal growth.

We collaborate.

We invite others to contribute to achieving our goals and value their time and expertise. We share what we know generously within our legislated limits, and seek opportunities to collaborate externally in areas of mutual interest.

We are innovators.

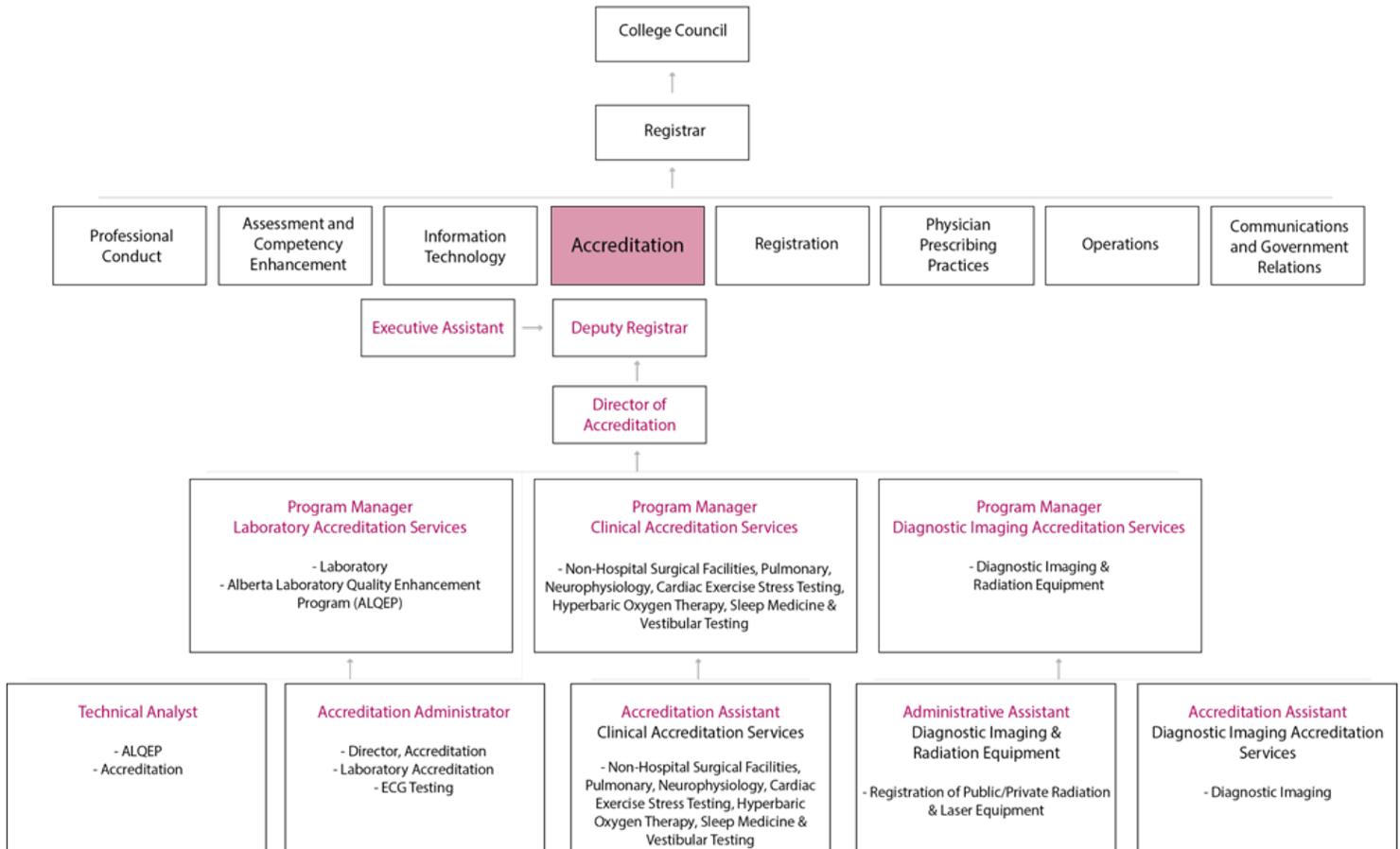
We think ahead to create opportunity. We set the bar high and value creativity in exploring new and better ways of doing our work.

We enjoy and find meaning in our work.

We care about what we do and give our best. While our work is serious, we enjoy camaraderie with our coworkers and take time to celebrate each other's milestones and achievements.

2.3 CPISA Organizational Structure - Figure 1

Accreditation Organizational Chart



2.4 Accreditation Program History

In 1965, the CPSA, upon recommendation from the Alberta Society of Pathologists, took steps to set up a program for accreditation for diagnostic medical laboratories. The Advisory Committee on Laboratory Medicine, which then reported to Council of the CPSA, was formed. The mandate of the Committee was to monitor and improve the quality of clinical laboratory services in Alberta. In order to meet this mandate, the Committee developed a process for accreditation that included requirements for on-site assessments of medical laboratories and a proposal for a proficiency-testing program to monitor testing performed.

The first assessments for accreditation took place in 1968 and included only non hospital-based laboratories. In 1970 the Alberta Department of Health entered into a contract with the CPSA to accredit hospital-based laboratories on their behalf and to make recommendations to them pertaining to accreditation.

The CPSA Accreditation scope has increased since then to include other public and/or private diagnostic programs in Alberta such as:

- Diagnostic Imaging
- Cardiac Exercise Stress Testing (CEST)
- Sleep Medicine
- Vestibular Testing
- Hyperbaric Oxygen Therapy (HBOT)
- Pulmonary Function Testing (PFT)
- Non-Hospital Surgical Facility (NHSF)

2.5 Authority and Oversight

The CPSA is constituted under the *Health Professions Act* (Schedule 21) with a mandate to regulate medical practitioners and medical practice in the best interests of the public of Alberta. Authority to accredit specified medical services and facilities is one aspect of that mandate.

Pursuant to section 8.4 of Schedule 21 of the *Health Professions Act*, and the Bylaws of the CPSA, DI facility staff are required to cooperate fully with any assessment, which shall include:

- a) permitting the assessment team to enter the imaging DI facility and assess the premises and all diagnostic equipment located therein;
- b) permitting the assessment team to assess all records pertaining to the provision of diagnostic imaging services, and providing copies of the same if so requested;
- c) providing to the assessment team, information requested by them in respect of the provision of diagnostic imaging services, in the DI facility;
- d) providing the information described in clause (c) in the form requested by the assessment team;
- e) providing requested samples or copies of any material, image, report, or product originating from the diagnostic imaging services, provided by the DI facility;
- f) answering questions posed by the assessment team as to procedures or standards of performance and if requested, providing copies of records relating to procedures followed and standards of performance applied in the diagnostic imaging DI facility;
- g) providing requested copies of all documents and information relating to business arrangements involving the practice conducted in the diagnostic imaging DI facility.

Although the CPSA's statutory authority does not extend to health services in approved hospitals or healthcare DI facilities operated by the Government of Canada or the Government of Alberta (*Health Professions Act* Schedule 21 - 8.1(1)), the value of practice uniformity between the private and public sectors and the credibility of the CPSA's programs have long been acknowledged by practitioners and government. Consequently, four of the CPSA's accreditation programs (laboratory medicine, diagnostic imaging, pulmonary function and neurophysiology) are under contract with government agencies to provide accreditation of public sector facilities.

The CPSA's accreditation programs are overseen by a standing committee, the Medical Facility Accreditation Committee (MFAC), with members appointed by the Council from diverse disciplines in clinical and diagnostic medicine. MFAC conducts a secondary review of practice standards developed by the accreditation advisory committees, hears argument on all changes to accreditation standards and reviews all facility accreditation and physician approval statuses. A member of the MFAC also attends a full meeting of the individual accreditation advisory committees each year and audits minutes of all other meetings to report on the diligence and objectivity of the work conducted.

The 5 standing advisory committees are composed of peer professionals (both physician/technical) who identify the needs and realities of Alberta stakeholders based on local practice.

2.6 Overview of Diagnostic Imaging (DI) Accreditation Program

The CPSA administers accreditation programs for those services that Council determines deserve explicit standards and verification of compliance with those standards, whether pertaining to the qualifications of physicians who provide them or the safety of those services to the public.

Accreditation looks at compliance, emphasizing continuous quality improvement and promoting optimum performance. More specifically, the CPSA's accreditation program looks closely at policies, processes and procedures to assess the safety and reliability of the service being provided, as well as the performance of the people involved and the product produced.

The DI Accreditation Program examines all aspects of imaging quality and operations including:

- organization, management and personnel
- quality management systems including policy, process and procedure
- physical DI facilities
- radiation equipment
- supplies, consumables
- imaging information systems and archival
- pre-examination, examination and post-examination activities
- quality assurance activities
- safety
- infection, prevention and control

The DI Accreditation Program is a peer review process with a goal to improve diagnostic imaging service provision and performance through objective evaluation. Assessors evaluate the DI facility's compliance with the specific requirements of a standard based on objective observation and assessment. All accreditation assessment findings are vetted by the Advisory Committee on Diagnostic Imaging to eliminate any potential personal assessor bias, ensure a consistent and thorough approach for all DI facilities, and to review standards for applicability to current best practice.

Benefits of CPSA DI Accreditation Program

- Assists DI facilities with the process of ensuring accuracy and reliability of imaging services
- Provides standards of practice and assesses compliance to the standards
- Identifies deficiencies that affect the quality of imaging services, as well as patient and staff safety
- Provides educational opportunities for both the DI facility being accredited and the assessment team
- Promotes uniformity in practice provincially – where variations in practice are counter-productive for the province.
- Promotes standardization and educational initiatives across Canada through interprovincial collaboration
- Maintains a comprehensive data repository for scope and complexity of imaging services, and resources within the province

- Promotes and ensures dialogue amongst imaging service providers and administrators on best practices and best ways to incorporate them into the workflow
- Encourages and facilitates peer review
- Ensures effective medical direction over medical practices so that business interests do not determine the standards of care

Confidentiality

All assessment findings are confidential and are only disclosed to parties explicitly associated with an assessment. Documented consent must be obtained from the assessed DI facility for release of assessment findings or accreditation certificates to other parties.

Frequency and Selection of DI facilities to be Assessed

DI facilities are assessed initially when opened, subsequently on a four year rotation and if they relocate their facility to a different physical location. This does not preclude an interim assessment that may be required as a result of expansion of imaging services or an unsatisfactory performance complaint.

Assessments are conducted by geographical zone areas ensuring that all DI facilities within the designated zone are assessed in the same calendar year. Ideally; the CPSA attempts to assess larger imaging group facilities at the same time. At the beginning of the year, all DI facilities in the area due to be assessed are identified by the CPSA, and the Assessment team is assigned. All DI facilities, both public and private, performing imaging examinations are required to undergo an assessment.

After a new DI facility is registered and initially accredited, it will then be added in to the regular zone geographical 4-year cycle. If the timing of this next 4-year cycle is very close to when the new DI facility was accredited, the CPSA may choose not to re-assess the DI facility.

On-going Self-Assessment

The CPSA DI accreditation process does not have a requirement for self-assessment. However, the DI General Standards require DI facilities to conduct formal internal audits of all system elements, both managerial and technical, at a frequency defined in their quality management system. The CPSA accreditation standard tools are a significant resource for self-audits as they promote a constant state-of-readiness. DI facilities are able to customize the standards tools by:

- documenting/embedding links to policies, processes, procedures, records, forms and labels beside the relevant standard
- utilizing the tool for the performance of comprehensive or targeted audits in between the 4-year assessments

2.7 Personnel

2.7.1 CPSA Diagnostic Imaging Accreditation Personnel and Roles

The Assistant Registrar for Accreditation has overall responsibility for the diagnostic imaging accreditation programs and is supported by the Director of Accreditation, the Program Manager for Diagnostic Imaging Services, the Accreditation Administrator, and the Administration Assistant for Radiation Equipment.

2.7.2 Advisory Committee on Diagnostic Imaging (ACDI)

The Advisory Committee on Diagnostic Imaging oversees the CPSA's accreditation program for medical diagnostic imaging DI facilities; for private DI facilities as defined in CPSA by-laws and for public DI facilities through contract with Alberta Health Services. Through the development of evidence based standards and monitoring DI facility compliance with those standards, the Committee promotes high standards of medical practice in diagnostic imaging facilities.

Roles and Responsibilities of the ACDI

- Develop and maintain evidence based standards for imaging practice
- Provide advice/recommendations to the Medical Facility Accreditation Committee (MFAC) on pending decisions relating to the provision of diagnostic imaging services
- Monitor compliance with CPSA approved standards through reviewing on-site assessment accreditation reports
- Provide education to promote safety and quality improvement initiatives
- Respond to the needs of stakeholders for improved imaging services in Alberta
- Review and audit of the business practices of the DI facility to ensure compliance with relevant CPSA By-laws and Standards

Membership and Tenure

Committee members are appointed by MFAC. Membership considers expertise, geographic location, urban versus rural and public versus private representation. Members, who serve by virtue of their position, serve as long as they fill that position.

Membership includes:

- Radiologists
- Cardiologists
- Diagnostic Imaging Technologist(s), Ultrasonographers
- CPSA staff (Assistant Registrar, Senior Medical Advisor, Director, Program Manager, Accreditation Assistant, Administrative Assistant, Radiation Equipment Assistant)

Non-Voting Members:

- Assessment Coordinators / CPSA Staff

All voting members are registered health professionals responsible to their respective professional regulatory body for their competence, their standards of practice and their conduct.

2.8 Assessment Teams

2.8.1 Assessment Coordinator

Each assessment team will include an Assessment Coordinator who is a consultant of the CPSA. Their primary role is to coordinate, organize and facilitate the assessment process.

2.8.2 Assessor (Quality Assessors and Modality Specific)

Selection of the assessment team is based on:

- scope and complexity of DI services
- number/geographic location of facilities
- experience of team members

The Assessment Coordinator(s) are present at each on-site assessment to promote consistency and continuity and to ensure an un-biased process.

2.8.3 Physician Reviewer

A Physician Reviewer will be assigned to an assessment team when:

- A new imaging facility has applied for Accreditation, and will require an initial image review
- An existing accredited imaging facility is applying for a new imaging service (modality), and will require an initial image review in the new imaging modality

2.8.4 Team Selection

The Assessment Coordinator(s), in collaboration with the CPSA, selects the members of the team which may include experienced imaging technologists, and imaging specialists (if required). All team members are provided with the training, information and material necessary to conduct a fair and thorough assessment.

Selection of the assessment team is based on:

- scope and complexity of imaging services
- number/geographic location of DI facilities
- experience of team members
- mitigation regarding conflict of interest

The Assessment Coordinator(s) are present at each on-site assessment to promote consistency and continuity and to ensure an un-biased process.

2.8.5 Assessment Team Training

All assessment team members are required to participate in a CPSA Assessor Training session, within 4-6 weeks prior to performing an on-site assessment. Following completion of the training session, assessment team members must demonstrate competency by successful performance of an on-line examination.

Upon successful completion of the training session and exam, all assessors receive a continuing professional development certificate.

2.8.6 Conflict of Interest / Confidentiality Agreements / Liability

All members of CPSA accreditation committees and assessment teams sign a Confidentiality agreement with the CPSA on an annual basis. Committee members and assessors are also required to confidentially destroy all confidential assessment materials or return to the CPSA for confidential disposal.

Assessment team members are also required to sign a Conflict of Interest agreement for each assessment cycle to ensure there are no potential conflicts specific to that assessment.

The CPSA's liability insurance specifically extends to cover assessors who are employed, contracted or act as agents. As well, the *Health Professions Act* extends liability protection to all CPSA staff, contractors and agents.

While performing assessments for the CPSA, assessors are advised not to display conduct that can be reasonably construed as a solicitation or offer consultant services that may compromise the objectivity of the assessment.

3.0 Standards Document

3.1 Standards Overview

The Standards are the basis for accreditation decisions and are compiled by CPSA and stakeholder experts, reviewed and are reviewed and approved by the Advisory Committee on Diagnostic Imaging, with final vetting and approval by the Medical DI Facility Accreditation Committee.

The Standards are evidence based and reference accepted best practices, Provincial and Canadian legislation, relevant International Organization for Standardization (ISO) standards, and other recognized provincial, national and international standards. Each accreditation standard has an accompanying reference citation(s).

All standards included in the documents are mandatory requirements for accreditation.

The Standards are process-based and incorporate a quality management system approach. The language, terms and organization of the documents are consistent with ISO 15189, where relevant.

A review of accreditation standards occurs on an ongoing basis, considering and incorporating stakeholder feedback. Comprehensive formal review occurs on an annual basis.

The CPSA Diagnostic Imaging Accreditation program currently maintains the following standards documents for the assessment of diagnostic imaging facilities:

- General (which includes Radiography, Fluoroscopy and Interventional)
- Bone Mineral Densitometry
- Computed Tomography
- Echocardiography
- Magnetic Resonance Imaging
- Mammography
- Nuclear Medicine and Positron Emission Tomography
- Ultrasound

All accredited Alberta DI facilities receive a complete standards document set. CPSA accredited imaging DI facilities and other approved users may access, print or make a copy of the standards for their non-commercial personal use. Any other reproduction in whole or in part requires written permission from the CPSA and the material must be credited to the CPSA.

Prior to each assessment, standards documents applicable to the scope of the imaging services of a DI facility will be made available to:

- DI facilities for self-assessment and/or to prepare for an on-site CPSA assessment
- CPSA assessors in preparation for on-site assessments and to record objective evidence/ observations while performing on-site assessments

3.2 Format of Standards

The standards are process-based and incorporate a quality management system approach. The language, terms and organization of the documents are consistent (where relevant) with ISO 15189/9001.

All standards documents are consistently organized in the following order (as applicable in each document):

- Organization, Management & Personnel
- Quality Management System
- Physical DI facilities
- Equipment, Consumables and Supplies
- Diagnostic Imaging Information Systems
- Pre-examination policies, processes and procedures
- Examination policies, processes and procedures
- Quality Assurance of examination procedures
- Post-examination policies, processes and procedures
- Safety
- Infection, Prevention and Control

The 'General Standards' document includes ALL standards common to ALL modalities. To eliminate redundancy, the modality-specific standards include ONLY those standards specific and relevant to each modality.

Figure 2 - Standard Document Format Example

#	Standard	Reference	Assessment of Compliance
IG.1.2 Organization & DI facility Management - Personnel			
IG.1.2.2	The DI facility has comprehensive written personnel comprehensive written policies, processes and procedures, which include comprehensive job descriptions that define qualifications and duties for all personnel.	Safety Code 35, Health Canada (A) AC ¹ RIS – 3.1 AC ² MIC - 5.1 CLSI QMS08-A2 – 6.3.1 IANZ ¹ - 5.1.1, 5.1.2, 5.1.7 ISO 15189 – 5.1.3, 5.1.9.a-d NCCLS ⁵ HS1-A2 – 5.3.1 RANZCR 4.1.1.2	Are there comprehensive written DI facility personnel (technical, medical and other) policies and job descriptions that define the qualifications, responsibilities and expectations of DI facility personnel?
			C <input type="checkbox"/> P <input type="checkbox"/> E <input type="checkbox"/> N <input type="checkbox"/> N/A <input type="checkbox"/> Observation:

Each standard consists of the following components:

- **CPSA standard number**
- **Patient or staff safety risk category** (where applicable):
 - Each standard has been reviewed to determine if it represents a direct and/or immediate patient or staff safety risk.
 - Those with either a patient safety (PS) or staff safety (SS) designation indicate that any non-compliance may have direct and/or immediate impact on safety.
 - PS/SS standards are ‘shaded’ for ease of detection
 - Assessors must ensure that ALL standards with either a PS or SS designation are directly assessed at the time of the on-site assessment.
- **Description of standard requirement**
- **Specific reference(s)** linked to reference listing at the end of the document
- **Interpretation guidance** where relevant regarding the application of requirements
- **Assessment of compliance questions (AOC)** that provide specific guidance and practical direction for evaluation of compliance with the standard
- **Compliance assessment category checkboxes**
- **Observation field for recording of objective evidence** (field is expandable in electronic document)

3.3 Assessment of Compliance (AOC)

- Although the AOC questions address the key evidence required to meet the intent of each standard, they **are not meant to be all encompassing**.
- There may be other evidence that demonstrates compliance with the intent of the standard. Individual assessors apply their own expertise in determining compliance with each standard.
- Compliance with the standard may be assessed by review of documents and records, observation, interviews or a combination of these techniques.
- Where AOCs state “All of the following”, compliance with all elements is expected to achieve compliance with the standard.

Assessment of Compliance Categories – the CPSA “PEN” or CPEN



Compliance Assessment Categories:	
C	meets intent and requirements of standard
P	in progress (working towards meeting intent and requirements of standard; assessor notes evidence of progress towards full compliance)
E	exceeds requirements of standard
N	does not meet intent and/or requirements of standard

N - Upon assessment of the objective evidence, failure to meet the intent and/or requirement of the standard will result in an assessment of non-compliance.

The standards are process based and a single non-compliance may encompass one or more observations. In assessing compliance with the standard, assessors will record direct specific objective evidence, which will be included in the report for each non-compliance.

P - “In Progress” citations require submission of future evidence of compliance based on direction from the assessor and/or the Advisory Committee. Examples where this assessment may be applied include situations such as: equipment purchased but not on-site and/or implemented; renovations in progress but not complete

Receipt of “FULL” accreditation status is contingent upon satisfactory resolution of all non-compliances (N and P).

E - “Exceeds Requirement” recognizes those situations where a DI facility exceeds the intent of the standard and employs commendable practice. The intent of capturing these occurrences is to promote and focus on quality initiatives.

3.4 Terms and Definitions

A listing of application terms and definitions is provided at the end of each standards document.

3.5 Reference Listing

A detailed reference listing is provided at the end of this document. Specific reference citation details can be accessed by clicking on individual link(s) included beside each standard. The references support the content and intent of each standard. It should be noted that all components of the cited references may not always be relevant and/or applicable. Compliance is expected with CPSA Standards; not the cited references.

3.6 Review and Revision of Standards

A comprehensive review of references occurs annually to ensure they are compliant with current standard references and best practices. Supporting references and any new references are reviewed, updated and their impact (if any) on the wording of the requirement is assessed.

Any stakeholder may offer suggestions for standards revision at any time.

Revision submissions are considered by the CPSA ONLY if they meet the following conditions:

- submitted using the Stakeholder Standards Review Form, found on the DI Accreditation website page
- identification of specific standard or section if applicable to multiple standards
- supported by detailed rationale/justification AND verifiable references (link or attachment must be included)
- applicable to all diagnostic DI facilities across the province and are not limited to organization specific practice
- contact information included for use by the CPSA if clarification of submission is required

4.0 Accreditation Process – New/Relocated DI facility Accreditation

		CPSA			Zone Group Facility	CPSA/Assessment Coordinator		Assessment team members / Physician Reviewers				
Initiation	Upon CPSA Notification	1	2	3	4							
Pre-Assessment	Time frames are dependent on the scheduling of the individual assessment	5	6	7	8	9						
	Differentiation between a relocation and a new DI facility or adding a new DI imaging service to an existing facility	10			11	12						
		13	14			15						
	Relocation: No phase	16			17	18	19					
	New facility – initiation and Phase I											
	New imaging service – initiation and Phase I											
~2-4 weeks prior to assessment	20	21					22					
~1-2 weeks prior to assessment	23											
On-site assessment	Day prior to assessment						24					
	Day of assessment				26	25	27	30	31	28	29	30
	End of assessment						32					
Post on-site	ASAP after each DI facility on-site assessment	34					33	35				
	Prior to Advisory Committee meeting / Meeting	36	37									
	Within 15 business days of Committee meeting	38	39									
	30/90 days past report distribution				40							
	Next Advisory Committee meeting post response receipt	42	43				41					
Stop here for Relocation. Continue on for New DI facility or New Imaging Service (existing facility)												
Phase I to Phase II (image review phase)	Submission of images to CPSA after 90 days of imaging	44/44(b)		46	47					45	46(b)	
	Next Advisory Committee meeting review	48	49	50	51	52	53	54				

4.1 Initiation – New/Relocated DI facility

Step	Responsibility	Task	Additional Information
1	CPSA Pre-assessment	Provides DI facility with assessment information New DI facility and New Modality (existing DI facility) is considered in Application - Phase I at this point. Relocations are not assigned a 'Phase'; as relocations do not require image submissions {Facility Application including Pre Assessment Data Verification Form} or {Notification of Change}	CPSA receives notification of: <ul style="list-style-type: none"> • New facility opening • New imaging service (modality) provision within existing facility • Facility relocation DI facility provides the CPSA with assessment information by filling out notification form for the new facility, new imaging service (modality) or relocation. DI facility provides name of accreditation contact. The CPSA sets up secure SharePoint access for the designated accreditation contact.
2	CPSA Pre-assessment	Selects AC and Physician Reviewer (if applicable)	Potential conflicts of interest are considered and mitigated when selecting proposed AC and Physician Reviewer (if applicable).
3	CPSA Pre-assessment	Provides DI facility to be assessed with the Assessment Logistics Form {ALF}	The zone area/group/DI facility Medical Director or authorized executive leadership designate is requested to complete and sign the Assessment Logistics Form (ALF) which includes: <ul style="list-style-type: none"> • provision of a key assessment /accreditation contacts for a zone/group/facility • approval of proposed Assessment Coordinator(s)
4	DI facility Pre-assessment	Completes Assessment Logistics Form	Completed form is submitted with signatures to CPSA within the specified timeline.

4.2 Pre-Assessment

Step	Responsibility	Task	Additional Information
5	CPSA Pre-assessment	Determines specific assessment dates in consultation with DI facility management {ALF}	CPSA liaises with the DI facility designated accreditation contact to determine specific assessment dates to facilitate the assessment prior to opening /provision of imaging services (as close as opening date as possible; turn-key).
6	CPSA Pre-assessment	CPSA liaises with DI facility accreditation contact regarding assessment process	CPSA provides the following information: <ul style="list-style-type: none"> • Ensure facility has program guide to review • Ensure facility has new DI Accreditation Standards • overview of assessment process steps • direction on the use of the Standards • assessment logistics & timelines DI facility training sessions are made available on the facility SharePoint and focus on: <ul style="list-style-type: none"> • overview of assessment process steps • assessment day expectations • use of the standards tool • assessment logistics & timelines

7	CPSA Pre-assessment	<p>Pre-assessment Data Verification Form completed along with the new facility or new imaging service form. {PADV}</p> <p>Relocations do not need to send a PADV in unless there is a change in the scope of imaging services</p>	<p>The PADV requests submission of the following for each individual DI facility undergoing assessment:</p> <ul style="list-style-type: none"> • general DI facility information • hours of operation • key DI facility personnel • scope of modalities (imaging services) to be provided or added • zone or centrally managed programs / processes • organizational structure • blank examples of facility imaging examination request (requisitions/consultation) forms and blank screening form/questionnaires) • List of all CPSA certified pieces of radiation equipment registered to the facility • Copy of the CAR-MAP certificate if providing diagnostic mammography imaging services • Copy of the CNSC certificate/licensure if providing nuclear medicine imaging services • Completed IG.5.0 Diagnostic Imaging Information Services Questionnaire <p>New DI facility / new imaging service (modality) registrations are requested to submit:</p> <ul style="list-style-type: none"> • organizational structure • examples of examination request (requisitions/consultation) forms and patient reports • zone/group/centrally managed programs / processes
8	DI facility Pre-assessment	<p>Completes Pre-assessment Data Verification Form and submits with required signature within the specified timeline {PADV}</p>	<p>CPSA reviews PADV for completeness.</p> <p>CPSA follows up directly with the DI facility accreditation contact regarding any missing documentation or documentation requiring further clarification.</p>
9	CPSA/AC Pre-assessment	<p>Selects assessment team members (assessors) based on zone/group/DI facility scope of modalities, availability and experience</p>	<p>The CPSA reviews the scope of modalities for each DI facility based on the submitted PADV documentation.</p> <p>The Assessment Coordinator does not assess; they are responsible for coordinating and leading the assessment team.</p> <p>Selection of the assessment team is based on:</p> <ul style="list-style-type: none"> • scope and complexity of imaging services • requirement for out-of-province assessors (if necessary) • number/geographic location of DI facilities • experience of team members • mitigation – conflict of interest (employment/affiliation) <p>CPSA ensures separate assessors (i.e. not the AC) are assigned for the assessment of ALL components of the General Standards.</p> <p>Each modality-specific assessor also reviews the relevant sections of the General Standards in conjunction with their modality specific assessment.</p>
10	CPSA Pre-assessment	<p>Advises DI facility Medical Director and/or</p>	<p>For each zone/group/DI facility (ies) assessment, the DI facility Medical Director and/or designated executive leadership receives a listing of</p>

		designated executive leadership of proposed assessment team members and requests formal written approval using the Proposed Team Member Form {Team Approval Form}	the proposed team members including their: <ul style="list-style-type: none"> • name • scope of assessment activities • location of employment/employer
11	DI facility Pre-assessment	Submits written approval of assessment team members to CPSA	If any original members are not approved by the zone/group/DI facility(ies) due to an identified conflict of interest, the CPSA will solicit alternate assessment team members and then request approval.
12	CPSA/AC Pre-assessment	Sends confirmation of team approval and assessment dates to assessors and ACs	CPSA sets up assessor access to Assessor SharePoint and sends notifications
13	CPSA Pre-assessment	Ensures relevant assessment tools (Standards documents) are available to identified DI facility accreditation contacts via Facility SharePoint	Facility SharePoint houses the General DI Standards, plus the relevant modality specific Standards available. There may be individual sections, as well as within sections, that are not applicable to each DI facility. Assessors will not be assessing these specific requirements.
14	CPSA Pre-assessment	Prepares documents for provision to Assessors	The CPSA prepares tools/supporting documentation for assessors: <ul style="list-style-type: none"> • PADV scope of modalities crosswalk for all DI facilities • modality specific assessor guides • New Facility/New Imaging service (modality) or Relocation Accreditation program guide
15	CPSA/AC Pre-assessment	Prepares final summary schedule and detailed DI facility schedules	Schedules are finalized. <ul style="list-style-type: none"> • if the assessment is a full day, then the facility will be asked to provide the lunch for the team via OLF; the CPSA reimburses the facility • If there are two assessments in one day, the morning facility will be asked to provide recommendations close for the team to obtain lunch in between facility assessments – via OLF
16	CPSA Pre-assessment	Distributes final summary and detailed DI facility schedules {Summary Assessment Schedule} {Individual Facility Schedule}	Final summary assessment schedules are uploaded to the DI facility and Assessor SharePoint sites. Detailed individual facility schedules are uploaded to the DI facility SharePoint site.
17	DI facility/CPSA Pre-assessment	Provides each DI facility to be assessed with an On-Site Logistics Form {OLF}	Coordinates assessment logistics with the DI facility/zone area accreditation contact. CPSA requests the following for zone/group/DI facility (ies) by requesting completion of the assessment On-site Logistics Form. Each DI facility is required to provide: <ul style="list-style-type: none"> • dedicated meeting room for the assessment team to work in • if the assessment is a full day, then the facility will be asked to provide the lunch for the team via OLF; the CPSA reimburses the

			<p>facility</p> <ul style="list-style-type: none"> • If there are two assessments in one day, the morning facility will be asked to provide recommendations close for the team to obtain lunch in between facility assessments – via OLF • access to any imaging records located outside the DI facility or department
18	CPSA/AC Pre-assessment	Sends completed On-site Assessment Logistics – Zone & DI facilities forms {OLF}	<p>Completed forms are submitted within the specified timelines.</p> <p>Zone/group/DI facility (ies) has identified key DI facility interviewees.</p> <p>CPSA confirms receipt of all information regarding the above arrangements and follows-up with DI facilities regarding any missing or conflicting information.</p>
19	CPSA/AC Pre-assessment	Coordinates team assessment logistics and sends to CPSA and team	<p>AC determines specific assessment team logistics to maximize efficiency, if required:</p> <ul style="list-style-type: none"> • transportation • accommodation • team meeting(s) • AC distributes assessment logistic details to CPSA and individual team members.
20	CPSA Pre-assessment	Distributes assessment documentation to each assessment team member to facilitate adequate preparation	<p>CPSA provides each team member the appropriate information for the assessment, including:</p> <ul style="list-style-type: none"> • completed PADV/PADV scope of modalities crosswalk • summary of previous citations and responses • modality specific assessor guides for each DI facility • copy of the General Standards and relevant Modality Standards • All assessment information is provided to team members via their secure SharePoint site
21	CPSA Pre-assessment	CPSA provides assessment team training session	<p>CPSA assessment team training: Mandatory for all assessment team members to participate in one session prior to the on-site assessment.</p> <p>Training is conducted by recorded webinar or on-line training module. Following completion of the training session, assessors must demonstrate competency by successful performance on an examination.</p> <p>Continuing education certificates are provided upon successful demonstration of competency</p> <p>Training sessions encompass:</p> <ul style="list-style-type: none"> • Overview of the CPSA assessment process and standards documents • General assessment guidance and techniques • CPSA assessor policies (e.g. confidentiality, conflict of interest, honoraria, expenses, etc.) • Specific assessment logistics
22	Assessment team Pre-assessment	Reviews assessment documentation and materials in preparation for the on-site assessment.	<p>Each assessment team member is expected to review the assessment documentation relevant to their scope of assessment activities to ensure that they are adequately prepared to perform a thorough and efficient assessment.</p> <p>The primary purpose is to:</p> <ul style="list-style-type: none"> • become familiar with the General and applicable modality-specific

			<p>standards</p> <ul style="list-style-type: none"> • become familiar with the scope of modality including which programs/processes are zone managed • identify areas of concern for further follow-up during the assessment (previous citations) <p>Standards tools can be customized by each assessor to meet their personal preferences for recording of observations and assessment categories on-site (e.g. add personal comments/directives, add additional space for recording, etc.)</p> <p>Assessors are expected to bring their own customized tools, either paper or electronic tablet version, to use during the assessment.</p>
23	CPSA Pre-Assessment	Provides assessment documentation to AC	CPSA prepares all relevant assessment documentation to the assessment team.

4.3 On-site assessment

Step	Responsibility	Task	Additional Information
24	AC/CPSA On-site	Conducts pre-assessment team briefing meeting	<p>Prior to the initiation of on-site assessment the AC(s) liaises with the team via email and disseminates information on:</p> <ul style="list-style-type: none"> • team introductions • assessment schedules and logistics review • CPSA assessor name tags distribution • discussion of areas for focus/concern • confirmation of areas of focus for each assessment team member • assessor questions/ clarifications
25	AC/CPSA On-site	Conduct an opening meeting with DI facility personnel	<p>At the beginning of the on-site assessment at each DI facility, the AC conducts an opening meeting for zone/group/DI facility(ies) personnel that encompasses:</p> <ul style="list-style-type: none"> • introductions • assessment logistics and timelines • assessment process outline • facility to determine if imaging specialist on site wishes to be interviewed
26	DI facility On-site	Conducts DI facility tour for assessment team members	An initial tour of the DI facility will give a general overview of the imaging operation and key personnel.
27	Assessment team members On-site	Conduct on-site assessments in areas of expertise	<p>The Assessment Process – General</p> <p>The accreditation assessment process involves:</p> <ul style="list-style-type: none"> • verifying compliance with the intent of accreditation standards • follow-up of previously identified areas of concern • interaction with imaging staff at all levels • review of zone managed areas (as identified by the facility) • imaging specialist on-site interview if the facility requests (AC to ask facility in the pre-assessment meeting; QMS assessor conducts requested interview) <p>Assessor Behavior</p> <ul style="list-style-type: none"> • engage in clear and concise dialogue with DI facility staff • explain the assessment process to DI facility staff, as required

			<ul style="list-style-type: none"> • exhibit positive body language • assess according to the standards (unbiased approach) • there are many ways to meet the intent of a standard • adopt an educational rather than a consultative OR punitive approach • the goal of the assessment is imaging improvement • do not act as a consultant • be conscious of timelines and assessment schedules and obligations • do not engage in confrontational behavior • be facilitative, professional • ask leading questions (“so explain to me how, or.. “where would you find...”) <p>The CPSA Assessment Tool The on-site assessment is performed using the DI facility specific standards document tools.</p> <p>Each assessor must utilize both the General Standards tool and the modality-specific Standards tool(s).</p> <p>The General Standards document includes ALL standards common to ALL modalities. To eliminate redundancy, the modality-specific standards include ONLY those standards specific and relevant to each discipline. For example, general quality control, equipment, supplies, consumables and procedure manual standards are not repeated in each discipline specific standard.</p> <p>Assessment of Compliance Although the AOC questions address the key evidence required to meet the intent of each standard, they are not meant to be all encompassing. There may be other evidence that demonstrates compliance with the standard. Where AOCs state “all of the following”, compliance with all elements is expected (e.g. test request form). Individual assessors apply their own expertise in determining compliance with each standard. Compliance with the standard may be assessed by review of documents and records, observation, interviews or a combination of these techniques.</p>
28	Assessment team members On-site	Conduct on-site assessments in areas of expertise CPSA Assessor Guides: <ul style="list-style-type: none"> • General (Rad, Fluoro, IR) • Bone Mineral Densitometry • Computed Tomography • Echocardiography • Mammography • Magnetic Resonance Imaging • Nuclear Medicine • Positron Emission 	<p>Guidance for Assessors</p> <p>When assessing DI facility imaging sections:</p> <ul style="list-style-type: none"> • It is not possible to review the entire scope of imaging operations <ul style="list-style-type: none"> ○ focus on areas of highest and lowest imaging volumes, likely problem areas and imaging results with highest impact on patient care ○ directly assess ALL standards with either a PS or SS designation ○ verify that all non-conformances cited on the previous assessment have been corrected ○ utilize the CPSA Assessor Guides to focus / direct the your on-site modality specific assessment • Review Zone/centrally managed programs / processes • Review documents (policies, processes and procedures - PPPs) and records <ul style="list-style-type: none"> ○ the assessor should choose a random, representative selection of documents, records, images and reports to review ○ assessors should not rely solely on documents, records, images

		<p>Tomography</p> <ul style="list-style-type: none"> • Ultrasound 	<p>and reports chosen or selected by the DI facility for review.</p> <ul style="list-style-type: none"> • Review PACS • Observe activities: <ul style="list-style-type: none"> ○ engage in meaningful dialogue with DI facility imaging and non-imaging staff (ask open ended questions such as: (what, when, where, why, who, how) ○ compare observed activities to the DI facility policies, processes and procedures ○ use techniques, such as: <ul style="list-style-type: none"> ▪ tracer method: follow a sample through pre-examination, examination & post-examination ▪ drill-down: further investigate areas of concern ▪ show/teach me: staff members describe a procedure as they perform it • Gather information: <ul style="list-style-type: none"> ○ always seek corroboration/validation/verification of findings ○ evaluate for significance ○ if the assessor determines (due to professional judgement) that the facility will require a Physician Reviewer intervention, they will start to collect examination information (accession numbers) of random examinations as per the 'minimum' number of exams 8.0). They will take this information to the AC; the AC will forward to the CPSA. When the report is discussed at the AC/DI, and the decision is to involve a PR, the site will be asked to submit the examinations that the assessor listed • Determine the scope and nature of potential citations: <ul style="list-style-type: none"> ○ is there a P/P/ or P? ○ is the P/P/ or P in compliance with the standards? ○ is the P/P/ or P being followed as written? ○ is there evidence of training/competency assessment for the activity? ○ is there acceptable documentation of the activity? ○ is the required review of the activity performed and documented? • Discuss / confirm potential deficiencies with DI facility representatives
<p>28</p>	<p>Assessment team members On-site</p>	<p>Conduct on-site assessments in areas of expertise</p>	<ul style="list-style-type: none"> • Record objective evidence: <ul style="list-style-type: none"> ○ as immediately as possible after encountering citation ○ using the assessment standards tool (paper or electronic) ○ do not rely on memory ○ be factual and thorough ○ provide ample background detail for interpretation and determination by the CPSA of the requirement/EOC • Photographic evidence for the Advisory Committee: <ul style="list-style-type: none"> ○ for safety related citations, consult with AC for necessity to corroborate observation with photographic evidence ○ AC will be responsible for notifying the DI facility contact and for taking required photographs ○ AC will ensure that no individuals or confidential information are identifiable in the photographs <p>Compliance Assessment Categories</p> <ul style="list-style-type: none"> • Non-conformances (N) <ul style="list-style-type: none"> ○ failure to meet the intent and/or requirement of the standard ○ The standards are process based and a single non-

			<p>compliance may encompass one or more observations.</p> <ul style="list-style-type: none"> • In-progress citations (P) <ul style="list-style-type: none"> ○ working towards meeting intent and requirements of standard; assessor notes evidence of timely progress towards full compliance ○ require submission of future evidence of compliance based on direction from the assessor and/or the Advisory Committee. ○ examples where this assessment may be applied include situations such as: equipment purchased but not on-site and/or implemented; renovations in progress but not complete ○ are not meant to address partial or incomplete compliance (e.g. incomplete manuals) • Exceeds requirement citations (E) <ul style="list-style-type: none"> ○ recognize those situations where a DI facility exceeds the intent of the standard and employs commendable practice. ○ the intent of capturing these occurrences is to promote and focus on quality initiatives.
29	Assessment team members On-site	Notify AC/CPSA immediately of any serious deficiencies that may have immediate impact on staff or patient safety	<p>Assessors encountering any situation that in their judgment, represents potential for significant immediate harm to staff or patients are directed to bring it to the attention of:</p> <p>#1 - the imaging personnel for immediate action as deemed appropriate</p> <p>#2 - AC who will consult with the CPSA immediately via telephone</p>
30	AC/Assessment team members On-site	Assessors communicate PEN findings to AC while on-site	<p>PEN Findings</p> <p>The AC will determine and communicate the timelines and frequency for debriefing assessors to obtain assessment PEN findings. At larger DI facility assessments this could be multiple times per day.</p> <p>AC will ask assessors to provide the following for each citation and record the details in the citation recording template:</p> <ul style="list-style-type: none"> ○ Standard number (if known) ○ Compliance assessment category (PEN) ○ Detailed observation/objective evidence ○ Comments (where applicable) <p>The AC ensures all citations include sufficient and clear detail in the objective evidence to facilitate the CPSA determination of the requirement, EOC, and timeline for EOC.</p> <p>The AC will reiterate back to the Assessors the information so that there is not a misunderstanding regarding the potential non-conformance, as well as ensuring there is enough detail given.</p> <p>If the assessor and/or AC is unable to determine the appropriate standard number to reference the citation, ACs are advised to record the other citation details and the CPSA will make the determination.</p>
31	AC On-site	Conduct pre-summation conference team meeting	<p>The AC de-briefs with the entire assessment team prior to the DI facility summation conference to determine and summarize key findings for presentation at the summation conference.</p>

			The AC will make particular note of systemic zone/group issues (e.g. document control in multiple DI facility imaging sections)
32	AC On-site	Conduct a summation conference for the zone/DI facility management and personnel	<p>The primary purpose of the summation conference is to highlight the key findings and outline the next steps in the assessment process.</p> <p>The AC serves as the primary spokespersons during the summation meeting in order to bring consistency of format and detail to the process.</p> <p>In person summation conferences are conducted at the DI facility at the end of the DI facility assessment.</p> <p>Summation conference agenda:</p> <ul style="list-style-type: none"> • Start by offering positive feedback on any processes or observations, examples and kudos to the facility • Acknowledgement of DI facility personnel for their cooperation and support of the accreditation process • Short review of the objectives of the accreditation process • Review of commendable findings and practices including any ‘E’ citations • Review of significant non-conformances. (The purpose of this is to ensure that there are no “significant surprises” in the report when received by the DI facility.) • Review of purpose and inclusion of interview findings in final reports • Overview of the next steps in the CPSA accreditation process including timelines for: <ul style="list-style-type: none"> ○ meeting of the ACDI to review the draft final report ○ distribution of final report ○ DI facility responses and submission of EOC • DI facility questions

4.4 Post-assessment

Step	Responsibility	Task	Additional Information
33	AC Post on-site	Verbally communicates overall findings from the on-site assessment to CPSA	<p>Immediately following the DI facility’s on-site assessment, the AC provides the CPSA with an overview of the findings from the assessment including any significant findings that would impact the DI facility receiving approval to commence testing and reporting of patient results.</p> <p>As soon as possible (same day, preferably) following each DI facility’s on-site assessment, the AC securely submits for each DI facility:</p> <ul style="list-style-type: none"> • Citation Recording Summaries including: <ul style="list-style-type: none"> ○ Standard number (if known) ○ Compliance assessment category (PEN) ○ Detailed observation/objective evidence ○ Comments (where applicable) ○ Completed imaging specialist interview, if applicable <p>AC should include any additional information or direction regarding on-site findings that would assist the CPSA in finalizing the requirements and requested EOC.</p>

<p>34</p>	<p>CPSA Post on-site</p>	<p>Communicates decision on commencement of imaging services to DI facility</p>	<p>Based on the communication received from the AC, the CPSA makes the determination on whether the DI facility can commence testing and reporting of patient results.</p> <p>The DI facility receives an immediate, formal electronic communication indicating:</p> <ul style="list-style-type: none"> • approval for commencement of imaging services <ul style="list-style-type: none"> ➤ Application Phase - Phase I for new DI facilities or new imaging service (modality) – PENDING ACDI review (committee can either support or revise this decision at the meeting) ➤ Full accreditation for relocated DI facility (continuation) <p>OR</p> <ul style="list-style-type: none"> • the requirement for resolution of identified significant non-conformances <ul style="list-style-type: none"> ➤ which would keep a facility from offering imaging services (whether new, or new modality (existing), or relocating) <p>Upon resolution of the identified significant non-conformances, the CPSA communicates the approval for commencement of imaging services PENDING ACDI review (committee can either support or revise this decision at the meeting)</p>
<p>35</p>	<p>AC Post on-site</p>	<p>Submits Citation Recording Summary to CPSA</p>	<p>As soon as possible (same day, preferably) following the DI facility's on-site assessment, the AC securely submits:</p> <ul style="list-style-type: none"> • Citation Recording Summary including: <ul style="list-style-type: none"> ○ Standard number (if known) ○ Compliance assessment category (PEN) ○ Detailed observation/objective evidence ○ Comments (where applicable) • Completed site interview forms where applicable <p>AC should include any additional information or direction regarding on-site findings that would assist the CPSA in finalizing the requirements and requested EOC.</p>
<p>36</p>	<p>CPSA Post on-site</p>	<p>Formats and finalizes draft DI facility report</p>	<p>Based on the Citation Recording Summary and the completed interview forms provided by the AC/ CPSA, the CPSA completes/finalizes the following for the DI facility report:</p> <ul style="list-style-type: none"> • DI facility demographics and key personnel • Assessment information and team details • Accreditation process date • DI facility Overview • Interview summaries (if applicable) • Citations: <ul style="list-style-type: none"> ○ Standard number ○ Safety Risk category ○ Compliance assessment category (PEN) ○ Detailed observation/objective evidence ○ Requirement ○ Evidence of Compliance (where applicable) ○ Timeline for submission of EOC

			<p>All requests for 90 day EOCs are based on the judgment of the assessors/CPSA/and the ACDI and include but are not limited to the following:</p> <ul style="list-style-type: none"> • All ‘P’ – ‘In Progress’ citations • Issues cited on previous assessment reports • All requirements categorized as PS/SS • Systemic/multi-DI facility issues <p>The CPSA ensures consistent/uniform:</p> <ul style="list-style-type: none"> • application of the standards based on similar observations • wording of requirements and EOC • timelines for submission of EOC
37	ACDI Post on-site	Vets and approves final draft DI facility report for new facilities and new modality (existing facility) and relocations	<p>At the next scheduled ACDI meeting, the AC for the assessment presents the report and represents the assessment team in responding to questions or clarification requests.</p> <p>At the meeting, the AC for the assessment presents the reports to the ACDI and represents the assessment team in responding to questions or clarification requests.</p> <p>ACDI reviews/revises/approves the DI facility assessment citations to:</p> <ul style="list-style-type: none"> • eliminate any personal bias • ensure consistent application of the standards from one assessor/assessment to another • endorse EOC requirement and timeline for EOC submission based on risk assessment • ensure standards/requirements reflect current best practice <p>If moving a facility from a Phase I to a Phase II:</p> <ul style="list-style-type: none"> • a request for images is included in the report 90 days post (done by CPSA) • The Reviewer must be available to review the images and submit a report within one week of the facility submission to the CPSA
38	CPSA Post on-site	Distributes final DI facility report to DI facility Medical Director and/or designated executive leadership	<p>Within 15-20 business days of the ACDI meeting, the CPSA posts the finalized individual DI facility report on the secure CPSA SharePoint site.</p> <p>The CPSA notifies the DI facility Accreditation contact or designated executive leadership that the final report is available electronically in the Facility SharePoint.</p> <p>The final report is formatted to include a section for a DI facility response to each individual non-conformance/in-progress citations.</p>
39	CPSA Post on-site	Provides accreditation evaluation forms to the DI facility and assessors	<p>To evaluate the effectiveness of the assessment process and customer satisfaction, the DI facility and the assessment team are asked to provide feedback on the Accreditation Evaluation Forms.</p> <p>Stakeholders are afforded the opportunity for anonymous comment.</p> <p>Results are compiled and reviewed annually by the CPSA and ACDI.</p> <p>Changes to process are implemented as appropriate based on feedback.</p>
40	DI facility	Submits a response to	DI facilities are required to electronically input their response directly

	Post on-site	requirements and requested evidence of compliance – if required	<p>into the report and embed any requested supporting documentation/EOC as applicable. Responses are uploaded by the designated accreditation contacts to secure DI facility SharePoint site.</p> <p>For requirements with requests for EOC: DI facilities must provide a response and required EOC based on timelines specified in the report (30 or 90 days from the date of the report).</p> <p>Responses to requirements without requests for EOC: DI facilities must provide a response within 90 days from the date of the report.</p>
41	CPSA Post On-site	Reviews zone/group/DI facility responses to requirements and requested evidence of compliance	<p>The CPSA reviews zone/group/DI facility responses to requirements and requested evidence of compliance and provides recommendations to the ACDI as to the appropriateness of the response.</p> <p>If required based on the DI facility responses, the CPSA will consult with the AC and/or the individual assessment team members for clarification.</p> <p>The CPSA provides a section in each report to facilitate the Committee responses.</p>
42	ACDI Post on-site	Reviews/approves DI facility responses	<p>At the next meeting following the receipt of 30 and 90 day responses, the ACDI reviews/revises/ approves the recommendations of the AC on the DI facility responses, with a view to:</p> <ul style="list-style-type: none"> • acceptability of response/corrective action • further action/clarification required <p>EOC may be required for any response based on submitted response, regardless of whether EOC was requested in the initial report.</p>
43	ACDI Post on-site	<p>Grants accreditation status:</p> <ul style="list-style-type: none"> • approval for commencement of imaging services <ul style="list-style-type: none"> ➤ Application Phase - Phase I to a Phase II for new DI facilities or new imaging service (modality) OR ➤ Full accreditation continued for relocated DI facility (continuation) 	<p>ACDI determines if any outstanding non-conformances (either due to volume or type of non-conformances) would substantiate a continuance of “Provisional” status.</p> <p><u>New facilities and new modality (existing facility):</u></p> <p><u>Application Phase - Phase I</u> Once the identified outstanding non-conformance(s) are satisfactorily addressed, the DI facility is moved from a Phase I to a Phase II accreditation status and can begin imaging services at this time</p> <p><u>Relocation</u> The DI facility can continue to offer the same imaging services (as before at the previous geographical location) at the <u>new</u> geographical location.</p> <p>Accreditation decisions of the ACDI are reviewed and approved by MFAC.</p> <p>If the ACDI recommends that a DI facility is denied accreditation, the DI facility may access the CPSA formal appeal process.</p>

Stop here for Relocation. Continue on for New DI facility or New Modality (existing facility)			
44 44(b)	CPSA/Assessor Phase II - image review	Communicate to DI facility the next steps regarding image submission	An Assessor will go on site and choose randomly the examinations that meet the 'Minimum' number of images after the facility has been imaging for 3 months.
45	DI facility Phase II - image review	Submits images and supporting documentation approximately 90 days after being moved from Phase I to Phase II	The Assessor will submit to the CPSA the examination list. The CPSA will contact the facility, and have the facility submit the examinations and appropriate supporting documentation within an appropriate timeframe.
46	CPSA Phase II - image review	Receives images and supporting documentation, and forwards them to Physician Reviewer – to be completed in two weeks' timeframe	The CPSA will receive the submission, and forward the submission to the designated Physician Reviewer.
46(b)	Physician Image Review	Physicians review images and send findings to the CPSA	The Physician Reviewer has two weeks to complete the review and submit results directly to the CPSA.
47	CPSA Phase II	CPSA collates physician review findings into a draft Phase II report and report is reviewed at next ACDI meeting	Phase II report is drafted which is the image review portion.
48	ACDI Phase II	Vets and approves final DI facility reports	At the meeting, the AC for the assessment presents the reports to the ACDI and represents the assessment team in responding to questions or clarification requests. ACDI reviews/revises/approves the DI facility assessment citations to: <ul style="list-style-type: none"> • Review any 30/90 day responses from the Phase I report • eliminate any personal bias • ensure consistent application of the standards from one assessor/assessment to another • endorse EOC requirement and timeline for EOC submission based on risk assessment • ensure standards/requirements reflect current best practice
49	CPSA Phase II	Distributes final DI facility report to DI facility Medical Director and/or designated executive leadership	Within 15-20 business days of the ACDI meeting, the CPSA posts the finalized individual DI facility report on the secure CPSA SharePoint site. The CPSA notifies the DI facility Accreditation contact or designated executive leadership that the final report is available electronically in the Facility SharePoint. The final report is formatted to include a section for a DI facility response to each individual non-conformance/in-progress citations.
50	DI facility Phase II	Submits a response to requirements and requested evidence of compliance – if required	DI facilities are required to electronically input their response directly into the report and embed any requested supporting documentation/EOC as applicable. Responses are uploaded by the designated accreditation contacts to secure DI facility SharePoint site. For requirements with requests for EOC: DI facilities must provide a response and required EOC based on

			<p>timelines specified in the report (30 or 90 days from the date of the report).</p> <p>Responses to requirements without requests for EOC: DI facilities must provide a response within 90 days from the date of the report.</p>
51	CPSA Phase II	Reviews zone/group/DI facility responses to requirements and requested evidence of compliance	<p>The CPSA reviews zone/group/DI facility responses to requirements and requested evidence of compliance and provides recommendations to the ACDI as to the appropriateness of the response.</p> <p>If required based on the DI facility responses, the CPSA will consult with the AC and/or the individual assessment team members for clarification.</p> <p>The CPSA provides a section in each report to facilitate the Committee responses.</p>
52	ACDI Phase II	Reviews/approves DI facility responses	<p>At the next meeting following the receipt of 30 and 90 day responses, the ACDI reviews/revises/ approves the recommendations of the AC on the DI facility responses, with a view to:</p> <ul style="list-style-type: none"> • acceptability of response/corrective action • further action/clarification required <p>EOC may be required for any response based on submitted response, regardless of whether EOC was requested in the initial report.</p>
53	ACDI Phase II	Recommends Full accreditation status	<p>ACDI determines if any outstanding non-conformances (either due to volume or type of non-conformances) would substantiate a reversion to “Provisional” status. If this decision is made, a “Provisional” certificate is issued and the DI facility is advised to replace their “Full” certificate with the “Provisional” certificate.</p> <p>Once the identified “provisional” non-conformance(s) are satisfactorily addressed, the DI facility is granted “Full Accreditation” status and a certificate is issued.</p> <p>Accreditation recommendations of the ACDI are reviewed and approved by MFAC.</p> <p>If the ACDI recommends that a DI facility is denied accreditation, the DI facility may access the CPSA formal appeal process.</p>
54	CPSA	Stakeholder/facility survey feedback	CPSA sends out assessment invoices as well as stakeholder survey

5.0 Honoraria and Expense Reimbursement

Expense Claims:

Assessors and assessment coordinators will be given pre-populated expense claim forms which include the name of the assessor/assessment coordinator.

The CPSA asks that meals and parking charges appearing on hotel receipts are separated out into appropriate categories on the claim form as these do not fall under the accommodation category.

Refer to the Honoraria and Expense Policy for additional guidance and information.

Honoraria Claims:

The CPSA provides an honorarium for:

- the CPSA Assessor Training session/Webinar
- travel time
- time spent on-site performing the assessment
- post-assessment follow-up with assessors – if required for confirmation of observations or for complex/specialty assessments

Assessment coordinators keep track of the breakdown of assessment team on-site assessment time, travel time and meals and submit this form to the CPSA for entry on the assessors' submitted claim forms. Assessors are advised to track their own time getting to and from the initial and final assessment sites.

Assessors and assessment coordinators will be given pre-populated honoraria claim forms which include the name of the assessor/assessment coordinator and the honoraria rate.

Within two weeks after the on-site assessment, expense and honoraria claim forms must be forwarded to the CPSA for processing for payment.

6.0 Assessment Fees

For Independent/Private DI facilities:

An invoice is issued after the ACDI has reviewed the report. Payment to the CPSA is due upon receipt of the invoice.

For Public DI facilities:

For all public DI facilities, Alberta Health Services is invoiced at the beginning of the year that the zone area is undergoing an assessment.

7.0 List of Accredited Diagnostic Imaging DI facilities

A current list of CPSA Diagnostic Imaging DI facilities is available on the CPSA website.

8.0 Examination/Image Review

CPSA Diagnostic Imaging facility Accreditation processes of exam/image review is not meant to assess individual imaging specialist competency; it is rather a benchmark of imaging quality services and processes.

Facilities will be required to submit images for review in the following circumstances:

- A new imaging facility is opening up
- A new imaging modality is being added to an existing imaging facility
- The Advisory Committee on Diagnostic Imaging (ACDI) reviews the Accreditation report for an on-site accreditation assessment and advises that an image review will be conducted based on review of the report evidence /findings, including (but not limited to) Standard IG.8.3:

<p>IG.8.3</p>	<p>The DI facility participates in a peer review / learning system for all imaging specialists.</p>	<p>CAR² RANZCR¹ – 5.6.1, 10.3.2.3</p>	<p>Does the facility participate in peer review / learning programs, appropriate to the scope and complexity of service?</p> <p>Is this process used in conjunction with the DI facility's quality program?</p> <p>Does this program assist in assessing the overall system to help the DI facility manage reporting accuracy and other quality improvements?</p> <p>Does the peer review / learning program include:</p> <ul style="list-style-type: none"> • reactive or proactive double reading with 2 physicians interpreting the same study? • random selection of studies to be reviewed on a regularly scheduled basis? • examinations and studies that are representative of each physician's specialty? • assessment of the agreement of the original report with subsequent review (or with surgical or pathologic findings)? <p>Is medical imaging specialist competence monitored by the collection of individual performance data by modality in the context of peer group comparison?</p> <p>Are there criteria and processes in place to assess the initial training/competency and on-going competency of imaging specialists?</p> <p>If an individual accountability review becomes necessary, is there a process that a new review is started that focuses on individual conduct, performance, mentorship, education and remediation?</p> <p>Does this process also encourage the DI facility to contact the appropriate regulatory body for advice where an accountability review is being conducted?</p>
----------------------	---	---	---

8.1 Public Imaging Facilities

The CPSA will liaise with the Zone to ensure appropriate Physician Reviewers (PR) have access to the electronic image portal/spoke, if required.

Assessors will randomly pick examinations on-site based on:

- Minimum number of examinations per modality as outlined by the CPSA
- Professional judgment, scope and complexity of the imaging facility provides
- Quality program in place for peer review program; or compatible alternate (IG. 8.3)

The DI facility will then be asked to upload the chosen examinations into the electronic portal. Same day upload is optimal. The DI facility will notify the CPSA when the images are ready for review. Examinations and associated reports/paperwork that can be taken away by the AC on the same day is optimal. Preference is for secured electronic submission of associated reports/paperwork, etc. The DI facility will have 5 business days from the date of the on-site assessment to provide the documentation.

The PR will have one business week to review the images and associated paperwork and submit findings back to the CPSA.

8.2 Private Imaging Facilities

Assessors will randomly pick examinations on-site based on:

- Minimum number of examinations per modality as outlined by the CPSA.
- Scope and complexity of the imaging facility provides

The DI facility will submit the examinations and the associated reports/paperwork to the CPSA via courier (one package) within five business days from the date of the on-site AC visit. Examinations/associated reports/paperwork that can be taken away by the assessor on the same day is optimal. Preference is for secured electronic submission of associated reports/paperwork, etc.

1. **Make and send two (2) copies of all studies (with the exception of Mammography).**
2. **Fill out the column “# of cases submitted” and include this form with the site submission.**
 - **Submit the amount of cases as outlined per modality.**
 - **Use the least amount of CDs/DVDs as possible.**
 - **Alternatively, images can be sent using one modality per USB thumb drive. (Echo studies shall be submitted on DVD's only due to the dynamic nature of the exam)**
 - **These must be in uncompressed DICOM format with a viewer included.**
 - **Divide your site submission into modalities.**
 - ✓ **Each modality must be submitted on a separate CD/DVD or USB thumb drive.**
 - ✓ **Do NOT divide within the modality**
 - **Supply all patient requisitions, reports and supporting documents in paper form.**
 - ✓ **Staple each patient's documents together.**
 - ✓ **Place these in the same order they appear on the viewer.**
 - **Before submitting, ensure that the CD/DVD/USB thumb drive can be opened and viewed on any computer.**

NOTE: If the CDs/DVD's and/or USB thumb drive of patient examinations are not submitted as instructed above, or are corrupt, the facility may be asked to resubmit

The CPSA will then courier the package and/or email the secure electronic documents to the PR. The PR will have one week to review the images and submit findings back to the CPSA.

8.3 Minimum Number of Examinations by Modality (for exam submission)

Minimum Number by Modality

Bone Densitometry

- 6 cases to be chosen, 3 must be follow up studies with the previous studies included
- The 3 follow up studies need to include the current study, most recent study and baseline study. If a patient has less than 3 studies, then just current and baseline are required

Computed Tomography

- Two head
- Two spine (cervical and Lumbar)
- One extremity (musculoskeletal)
- 1 chest
- 1 abd/pelvis
- One chest/abdomen/pelvis

Echocardiography

- 4 cases to be chosen (2 normal cases and 2 significantly abnormal cases)
- If transeophageal, stress and contrast are performed, 3 additional (one of each)
- Full TTE with moving images, including 2D in all standard views/windows, and spectral Doppler and M-mode

Fluoroscopy

- One GI tract – lower
- One GI tract – upper
- One GU tract – hysterosalpingogram
- One GU tract – Cystogram
- One spine
- Two pain management

Mammography – Canadian Association of Radiologists-Mammography Accreditation Program (CAR-MAP)

No image / report submission required. Assessor will review random exams and images on site for technical review.

Mammography – Non CAR-MAP

Assessor will review random exams and images on site for technical review.

- biopsy
- screening

Magnetic Resonance Imaging

- Two head
- Two spine (cervical and lumbar)
- Two musculoskeletal
- 2 cardiac
- 1 abdomen/pelvis
- 1 chest
- 1 magnetic resonance angiography
- 2 pediatric examinations

Positron Emission Tomography/PET-CT

- Two FDG PET/CT

Nuclear Medicine

- One bone (3 phase)
- One whole body bone
- One lung
- One renal

- One HIDA
- One cardiac
- One liver/spleen
- One pediatric examination

Radiography

- Two abdomen/pelvis (one of female child bearing age)
- Two Pelvic girdle
- Two lower extremity
- Two upper extremity
- Two spine
- Two chest
- Two mobiles

Ultrasound

- Two Abdomen/Pelvis
- 2 Interventional
- 6 Obstetrical (one of the two is a trans-vaginal study, if performed, and of the four – two from each trimester)
- 1 Obstetrical Doppler
- 2 Gynecological
- 2 Vascular
- 1 Small Parts
- 1 MSK