

Schedule of Facility Fees

For Independent Health Facilities

Under the *Independent Health Facilities Act*

GENERAL PREAMBLE

1. Every licensee is responsible for ensuring that facility fees are charged to the Ministry, and payment accepted, only in accordance with the *Independent Health Facilities Act* (IHFA) and its regulations.
2. Facility fees shall be charged to the Ministry only in respect of a service rendered by a physician for which an amount payable is prescribed by the regulations under the *Health Insurance Act* (HIA), or a service prescribed as an insured service under the HIA rendered by a practitioner within the meaning of that Act (i.e., OHIP-insured medically necessary services provided to an insured person pursuant to a requisition.*)
3. Previous payment of a facility fee shall not be construed as approval of any particular billing practice.
4. Each independent health facility (IHF) licence is issued with respect to a specified single location or, in the case of mobile IHFs, with respect to specified multiple locations. Licensees are not permitted to charge facility fees to the Ministry, or to receive payment, in respect of services provided at locations other than the location(s) specified on the IHF licence. The unique billing number issued by the Ministry to each IHF shall only be used to charge facility fees to the Ministry for services provided at the location(s) specified on the facility licence. Non-compliance may lead to recovery of funds, licensing action in accordance with the IHFA, prosecution pursuant to the *Provincial Offences Act*, and/or such other legal action may be appropriate in the circumstances.
5. Where a referring physician requests a single site imaging study (for example, one breast, one limb), any additional imaging of a portion of the anatomy for comparison purposes is not an insured service and shall not be charged to the ministry.
6. Where a referring physician requests a single site imaging study, any additional imaging study is not an insured service and shall not be charged to the ministry unless the additional study is medically necessary as requested by the radiologist or referring physician and documented in the patient's record.
7. Where a licensee provides breast ultrasound services, a scan of the axilla is an integral part of the breast imaging exam. The licensee shall not charge any facility fees to the ministry in connection with an additional insured service fee code such as J182 (extremity ultrasound).
8. Where a referring physician requests mammography, the addition of ultrasound breast imaging services shall not be charged to the ministry unless the additional study is medically necessary as requested by the radiologist or referring physician and documented in the patient's record.
9. Where a copy of an imaging study is requested for the purpose of continuing medical care, the licensee shall not charge any person for costs of preparing a CD or other imaging media. If a licensee charges a patient in such circumstances, the ministry will reimburse the patient and recover the full amount from the licensee through set-off from future billings, in addition to applying an administrative penalty of \$50 per occurrence, pursuant to the IHFA and regulations.

* "written requisition" means: a written requisition from a referring physician or a requisition from a practitioner as may be permitted under the IHFA or the HIA and the regulations.

NUCLEAR MEDICINE – IN VIVO

PREAMBLE

SPECIFIC ELEMENTS

For Facility Fee Component (F fee)

- A. Preparing the patient for the procedure.
- B. Performing the diagnostic procedure(s).
- C. Making arrangements for any appropriate follow-up care.
- D. Providing records of the results of the procedure to the interpreting physician.
- E. Discussion with, and providing information and advice to, the patient or patient's representative, whether by telephone or otherwise, on matters related to the service.
- F. Preparing and transmitting a written, signed and dated interpretive report of the procedure to the referring physician.
- G. Providing premises, equipment, supplies and personnel for all *specific elements* of the facility fee components.

OTHER TERMS AND DEFINITIONS

1. Professional and facility fee components are claimed separately. Claims for the facility fee component F are submitted using listed fee code with suffix B. Where the IHF is submitting professional fee claims on behalf of the interpreting physician, claims for professional component are submitted using fee code with suffix C (e.g. J802C).
2. If examination of Brain, Lung, Liver or Spleen is limited to one view, the benefit (F fee) is to be reduced by 50%.
3. Repeat studies on the same day may be claimed only after exercise or drug intervention.

NUCLEAR MEDICINE – IN VIVO

Code	F
<u>CARDIOVASCULAR SYSTEM</u>	
Venography	
J802	99.75
- peripheral and superior vena cava	
First Transit	
J804	16.65
J867	59.35
- without blood pool images	
- with blood pool images	
Cardioangiography	
J806	98.45
- first pass for shunt detection, cardiac output and transit studies	
Myocardial Perfusion Scintigraphy	
J807	225.25
J866	45.05
J808	82.95
J809	45.05
- resting, immediate post stress	
- application of SPECT (maximum one per examination), to J807	
- delayed	
- application of SPECT (maximum two per examination), to J808	
Myocardial scintigraphy	
J810	91.35
- acute infarction, injury	
Myocardial wall motion	
J811	98.45
J812	49.85
J813	139.95
J814	49.85
- studies	
- repeat same day (to a maximum of three repeats)	
- studies with ejection fraction	
- repeat same day (to a maximum of three repeats)	
Note: J811 and/or J812 rendered in conjunction with J813 and/or J814 are insured services payable at nil.	
J815	136.35
Detection of venous thrombosis using radioiodinated fibrinogen up to ten days	
<u>ENDOCRINE SYSTEM</u>	
Adrenal scintigraphy	
J816	399.55
J868	467.30
J869	575.00
- with iodocholesterol	
- with iodocholesterol and dexamethasone suppression	
- with MIBG	
Thyroid scintigraphy	
J818	66.40
J871	106.75
- with Tc99m or I-131	
- with I-123	
Thyroid	
J817	29.65
J870	15.15
- uptake	
- repeat	

NUCLEAR MEDICINE – IN VIVO

Code		F
ENDOCRINE SYSTEM (continued)		
Parathyroid scintigraphy		
J820	-dual isotope technique with T1201 and Tc99m Iodine	243.00
J872	Metastatic survey with I-131	249.10
GASTROINTESTINAL SYSTEM		
Schilling test		
J821	- single isotope	46.25
J823	- dual isotope	49.85
Malabsorption test		
J824	- with C ¹⁴ substrate	59.35
J873	- with whole body counting	142.55
Gastrointestinal		
J825	- protein loss	85.35
J874	- blood loss using – Cr ⁵¹	64.10
J829	- transit	106.75
Calcium absorption		
J826	- Ca ⁴⁵	64.10
J875	- Calcium ⁴⁷ absorption/excretion	262.05
J827	Oesophageal motility studies – one or more	123.10
Gastro-oesophageal		
J876	- reflux	58.70
J877	- aspiration	41.55
Abdominal scintigraphy – for gastrointestinal bleed		
J830	- Tc99m sulphur colloid or Tc ⁰⁴	90.10
J878	- labelled RBCs	148.25
J879	- LeVeen shunt patency	68.65
J831	Biliary scintigraphy	118.55
J832	Liver/spleen scintigraphy	82.95
J833	Salivary gland scintigraphy	99.65
GENITOURINARY SYSTEM		
J834	Dynamic renal imaging	99.65
Computer assessed renal function		
J835	- includes first transit	136.35
J880	- repeat after pharmacological intervention	46.45

NUCLEAR MEDICINE – IN VIVO

Code		F
GENITOURINARY SYSTEM (continued)		
J836	Static renal scintigraphy	34.45
J837	ERPF by blood sample method	41.55
J838	GFR by blood sample method	41.55
J839	Cystography for vesicoureteric reflux	124.80
Testicular and scrotal scintigraphy		
J840	- includes first transit	85.35
HAEMATOPOIETIC SYSTEM		
J841	Plasma volume	45.05
J843	Red cell volume	49.85
J847	Ferrokinesis – clearance, turnover, and utilization	415.15
J848	Red cell, white cell or platelet survival	106.25
J849	Red cell survival with serial surface counts	153.50
Bone marrow scintigraphy		
J881	- whole body	117.70
J882	- single site	87.85
In-111 leukocyte scintigraphy		
J883	- whole body	377.20
J884	- single site	332.15
MUSCULOSKELETAL SYSTEM		
Bone scintigraphy		
J850	- general survey	107.35
J851	- single site	87.85
Gallium scintigraphy		
J852	- general survey	183.85
J853	- single survey	128.10
Application of Tomography (SPECT)		
J819	- where each SPECT image represents a different organ or body area, to J852, maximum three images per examination add	45.05
<p>Note: J850 and J851 are not to be billed together. J804 may be claimed in addition to J850 or J851 for blood pool study.</p>		
NERVOUS SYSTEM		
CSF circulation		
J857	- with Tc99m or I-131 HAS	124.50
J885	- with In-111	319.10

NUCLEAR MEDICINE – IN VIVO

Code		F
NERVOUS SYSTEM (continued)		
J886	- via shunt puncture	91.70
J858	Brain scintigraphy	93.60
RESPIRATORY SYSTEM		
J859	Perfusion lung scintigraphy	88.95
J887	Ventilation lung scintigraphy	111.50
J860	Perfusion and ventilation scintigraphy – same day	177.95
MISCELLANEOUS		
J861	Radionuclide lymphangiogram	116.15
J862	Ocular tumour localization	78.30
J864	Tear duct scintigraphy	100.80
J865	Total body counting	194.60
Application of Tomography (SPECT), other than to J808 or J852		
J866	- maximum one per Nuclear Medicine examination	add 45.05
SCINTIMAMMOGRAPHY		
<p>Scintimammography is not eligible for payment unless at least one of the following conditions is met:</p> <ol style="list-style-type: none"> a. the patient has a dense breast(s) and one or both of the following risk factors: <ol style="list-style-type: none"> i. a first degree relative with breast cancer diagnosed prior to age 50; or ii. a first degree relative with breast cancer diagnosed over age 50 and patient is within 5 years of the age when the relative was diagnosed with breast cancer. b. architectural distortion of the breasts due to prior breast surgery, radiotherapy, chemotherapy or the presence of breast prosthesis rendering mammography interpretation difficult; c. malignant breast lesion when mammography is unable to exclude multifocal disease; or d. solitary lesion identified on mammography of greater than 1 cm 		
Scintimammography		
J863	- unilateral or bilateral	103.50
<p>Note: For the purpose of this provision, “dense breast(s)” means (a) breast(s) occupied by over 75% fibroglandular tissue as noted on mammography.</p>		

DIAGNOSTIC RADIOLOGY

PREAMBLE

SPECIFIC ELEMENTS

For Facility Fee Component (F fee)

- A. Preparing the patient for the procedure.
- B. Performing the diagnostic procedure or assisting in the performance of fluoroscopy.
- C. Making arrangements for any appropriate follow-up care.
- D. Providing records of the results of the procedure to the interpreting physician.
- E. Discussion with, and providing information and advice to, the patient or patient's representative, whether by telephone or otherwise, on matters related to the service.
- F. Preparing and transmitting a written, signed and dated interpretive report of the procedure to the referring physician.
- G. Providing premises, equipment, supplies and personnel for all *specific elements* of the technical components.

OTHER TERMS AND DEFINITIONS

1. Professional and facility fee components are claimed separately. Claims for facility fee component F are submitted using the listed fee code with suffix B.
2. If less than the minimum number of views are performed, reduce listed fees by 25%.
3. If insured diagnostic radiology procedures yield abnormal findings or if they would yield information which in the opinion of the radiologist would be insufficient governed by the needs of the patient and the requirements of the referring physician or practitioner, the radiologist may add further views and claim for the additions which are to be noted in the report.
4. Where a referring physician requests a single site imaging study (for example, one breast, one limb), any additional imaging of a portion of the anatomy for comparison purposes is not an insured service and shall not be charged to the ministry.
5. A stereo pair is to be counted as two views.
6. No additional claim is warranted for the use of the image intensifier in diagnostic radiology.
7. Nasal bones or accessory nasal sinuses should not be routinely claimed in skull examination requests.

DIAGNOSTIC RADIOLOGY

8. Mandible X006 and Temporomandibular joints X007 are not both to be routinely claimed on the same patient but only when specifically ordered.
9. Conventional films of the spine should not be routinely done and claimed for before myelography. The necessity of having plain film studies of the spine prior to interpreting the myelographic studies is obvious. It is not essential, however, that these be done at the institution where the myelogram was done. If they have been done at an outside office, then it is a matter for the radiologist and the referring physician to have the films available. If they cannot be made available to the radiologist, it is an acceptable practice for him to do the required procedure of these areas and to claim for them so that they may be available for interpretation along with the myelographic study.
10. Lumbar or lumbosacral spine X028 does not include the entire sacrum. An x-ray of the sacrum may be carried out and claimed for only when specifically indicated.
11. Three or more views of the chest should not be done routinely and claimed when a chest examination is requested.
12. Chest studies should not be routinely done and claimed in mammography cases.
13. Fluoroscopy claims should not be submitted for any examination performed by the radiologist where fluoroscopy is generally regarded as an integral part of the examinations, e.g. examinations of the GI tract, urinary tract, and special procedures.
14. 'Colon - air contrast' may be claimed when performed according to generally accepted criteria. The colon should be scrupulously prepared. Five to eight full size views of the abdomen should be obtained after fluoroscopically controlled introduction of air and barium.
15. 'Oesophagus, stomach and duodenum - double contrast' presupposes the introduction of gas, the use of antifoam agent and a suitable barium mixture.
16. 'Pharynx and oesophagus - cine or videotape' (X106) should not be claimed routinely with X108 and X109 but only when specifically indicated.
17. Abdomen and chest studies should not be routinely done and claimed in gastrointestinal examinations.
18. Abdomen and/or pelvis should not be routinely claimed in lumbar spine examination requests.
19. A survey film of the abdomen is a single view. The ordering of additional films should be left to the discretion of the radiologist who has the authority to determine what examination is adequate for a specific patient. Obviously, if progress of a long tube is being followed, a survey film is sufficient. If, however, an intestinal obstruction is being followed, a single film is usually inadequate.
20. No extra fee should be claimed for rapid sequence IVP.
21. Nephrotomography is covered by the listings for intravenous pyelogram and planigram.

DIAGNOSTIC RADIOLOGY

22. For the following surgical procedures, the technical fee components for chest x-ray, X090, X091, X092 are only eligible for payment in the preoperative preparation of the patient when the referring physician obtains prior authorization of payment from the Ministry of Health and demonstrates the medical necessity of the service:

- a. cataract surgery;
- b. colonoscopy;
- c. cystoscopy;
- d. carpal tunnel surgery; or
- e. arthroscopic surgery.

[Commentary:

If there is an indication requiring a chest x-ray other than strictly for preoperative preparation for the above surgical procedures, prior approval is not required and the technical fee may be claimed.]

23. Mammography or x-ray of the chest, ribs, arm, wrist, hand, leg, ankle or foot, rendered in an Independent Health Facility or a hospital in-patient or out-patient department is insured in accordance with the Health Insurance Act when referred by a registered nurse holding an extended certificate of registration (RN(EC)).

DIAGNOSTIC RADIOLOGY

Code	F
HEAD AND NECK	
Skull	
X001	33.35
X009	41.55
X003	16.60
Facial bones	
X004	24.20
Nose	
X005	16.60
Mandible	
X006	24.20
X012	33.35
X007	24.20
Mastoids	
X010	31.95
X011	24.20
Note: Dental x-rays of the teeth are not an insured benefit.	
X016	16.55
X017	17.05
X018	18.80
X019	15.35
Neck for soft tissues	
X020	15.35
SPINE AND PELVIS	
Cervical spine	
X025	28.90
X202	37.25
X203	45.00
Thoracic spine	
X027	26.35
X204	33.35

DIAGNOSTIC RADIOLOGY

Code	F
SPINE AND PELVIS (continued)	
Lumbar or lumbosacral spine	
X028	28.90
X205	37.25
X206	45.00
Entire spine (scoliosis series)	
X032	59.70
X033	24.20
X031	33.10
Sacrum and/or coccyx	
X034	26.70
X207	34.60
Sacro-iliac joints	
X035	24.20
X208	32.30
Pelvis and/or hip(s)	
X036	16.60
X037	30.95
X038	35.55
UPPER EXTREMITIES	
Clavicle	
X045	16.60
X209	25.55
Acromioclavicular joints (bilateral) with or without weighted distraction	
X046	24.20
X210	33.00
Sternoclavicular joints (bilateral)	
X047	20.00
X211	28.55
Shoulder	
X048	20.00
X212	28.55
Scapula	
X049	20.00
X213	28.75

DIAGNOSTIC RADIOLOGY

Code		F
UPPER EXTREMITIES (continued)		
Humerus including one joint		
X050	- two views	16.60
X214	- three or more views	25.35
Elbow		
X051	- two views	16.60
X215	- three or four views	25.55
X216	- five or more views	34.40
Forearm including one joint		
X052	- two views	16.60
X217	- three or more views	25.55
Wrist		
X053	- two or three views	16.60
X218	- four or more views	25.55
Hand		
X054	- two or three views	16.60
X219	- four or more views	25.55
Wrist and hand		
X055	- two or three views	24.20
X220	- four or more views	30.85
Finger or thumb		
X056	- two views	12.80
X221	- three or more views	16.60
LOWER EXTREMITIES		
Hip (unilateral)		
X060	- two or more views	26.50
Femur including one joint		
X063	- two views	16.60
X223	- three or more views	24.75
Knee including patella		
X065	- two views	16.60
X224	- three or four views	25.55
X225	- five or more views	34.40
Tibia and fibula including one joint		
X066	- two views	16.60
X226	- three or more views	25.55

DIAGNOSTIC RADIOLOGY

Code		F
<u>LOWER EXTREMITIES (continued)</u>		
Ankle		
X067	- two or three views	16.60
X227	- four or more views	25.55
Calcaneus		
X068	- two views	16.60
X228	- three or more views	25.55
Foot		
X069	- two or three views	16.60
X229	- four or more views	25.55
Toe		
X072	- two views	12.80
X230	- three or more views	16.60
X064	Leg length studies (orthoroentgenogram)	24.20
<u>SKELETAL SURVEYS</u>		
Skeletal survey for bone age		
X057	- single film	16.60
X058	- two or more films or views	24.20
Other survey studies – e.g. rheumatoid, metabolic or metastatic		
X080	- single view	8.30
X081	- each additional film or view	8.30
<u>CHEST AND ABDOMEN</u>		
Chest		
X090	- single view	16.60
X091	- two views	24.40
X092	- three or more views	31.40
Note: Miniature chest film for survey purposes only is not an insured benefit.		
Ribs		
X039	- two or more view	20.00
Sternum		
X040	- two or more view	20.00
Thoracic inlet		
X096	- two or more view	16.60

DIAGNOSTIC RADIOLOGY

Code		F
<u>CHEST AND ABDOMEN (continued)</u>		
Abdomen		
X100	- single view	16.60
X101	- two or more views	25.40
<u>GASTROINTESTINAL TRACT</u>		
Palatopharyngeal analysis		
X105	- cine or videotape	32.90
Pharynx and oesophagus		
X106	- cine or videotape	32.90
X107	Oesophagus when X103, X104, X108 or X109 not claimed	29.75
Oesophagus, stomach and duodenum		
X108	- including survey film, if taken	51.65
X104	- double contrast, including survey film, if taken	54.10
X103	- double contrast, including survey film, if taken, and small bowel	67.95
X110	Hypotonic duodenogram	43.90
X109	Oesophagus, stomach and small bowel	65.90
Small bowel only		
X111	- when only examination performed during patient's visit	29.45
Colon		
X112	- barium enema including survey film, if taken	53.95
X113	- air contrast, primary or secondary, including survey films, if taken	68.35
Gallbladder		
X114	- one or multiple day examinations	33.40
X120	- one or multiple day examinations with preliminary plain film	44.40
X116	T-tube cholangiogram	24.20
X123	Operative pancreatogram or ERCP	24.20
<u>GENITOURINARY TRACT</u>		
X129	Retrograde pyelogram, unilateral or bilateral	24.20
X130	Intravenous pyelogram including preliminary film	55.35
X137	Cystogram (catheter)	26.60
X135	Cystourethrogram, stress or voiding (catheter)	30.65
X131	Cystourethrogram (non-catheter)	6.40
X191	Intestinal conduit examination or nephrostogram	24.20
X138	Percutaneous antegrade pyelogram	24.20
X139	Percutaneous nephrostogram	24.20

DIAGNOSTIC RADIOLOGY

Code		F
GENITOURINARY TRACT (continued)		
X134	Retrograde urethrogram	20.00
X136	Vasogram	20.00
X141	Cavernosography	23.05
OBSTETRICS AND GYNAECOLOGY		
X147	Hysterosalpingogram	33.25
FLUOROSCOPY – BY PHYSICIAN WITH OR WITHOUT SPOT FILMS		
X195	Chest	10.30
X196	Skeleton	10.30
X197	Abdomen	10.30
X189	Fluoroscopic control of clinical procedures done by another physician per ¼ hour	8.15
SPECIAL EXAMINATIONS		
Abdominal, thoracic, cervical or cranial angiogram by catheterization		
	Using single films	
X179	- non-selective	33.00
X180	- selective (per vessel, to a maximum of 4)	43.40
	Using film changer, cine or multiformat camera	
X181	- non-selective	66.50
X182	- selective (per vessel, to a maximum of 4)	88.40
X140	- selective (5 or more vessels)	353.85
Carotid angiogram by direct puncture		
X160	- unilateral	54.50
X161	- bilateral	87.65
Peripheral angiogram		
X174	- unilateral	33.25
X175	- bilateral	43.90
X198	Splenoportogram	65.90
X199	Translumbar aortogram	65.90
Vertebral angiogram – direct puncture or retrograde brachial injection		
X132	- unilateral	54.50
X133	- bilateral	89.10
X156	Arthrogram, tenogram or bursogram	29.25
X200	- with fluoroscopy and complete positioning throughout by physician	40.90

DIAGNOSTIC RADIOLOGY

Code	F
SPECIAL EXAMINATIONS (continued)	
Bronchogram	
X158	32.30
X159	42.80
X122	32.90
BONE MINERAL DENSITY (BMD) MEASUREMENT	
Dual-energy X-Ray Absorptiometry (DXA) – by axial technique only	
<p>Definition: For the purpose of second and subsequent testing,</p> <p>“high risk patient” means a patient;</p> <ol style="list-style-type: none"> 1. at risk for accelerated bone loss (in the absence of other risk factors, patient age is deemed not to place a patient at high risk for accelerated bone loss); 2. with osteopenia or osteoporosis on any previous BMD testing; or 3. with bone loss in excess of 1% per year as demonstrated by previous BMD testing. <p>“low risk patient” means a patient who is not a high risk patient</p> <p>Definition/Required Elements of Service: BMD measurement by DXA is an insured service only when all the following conditions have been met:</p> <ol style="list-style-type: none"> 1. the service is rendered for the prevention and management of osteoporosis or osteopenia; 2. when more than one site is measured, the sites include both hip and spine and where measurement of both hip and spine is not technically feasible the site measured consists of either hip or spine. <p>[Commentary: Measurement of hip and spine would be considered not technically feasible due to prosthesis or deformity.]</p>	
Baseline Test	
X145	47.75
X146	61.55
Second test - low risk patient	
X152	47.75
X153	61.55
Subsequent test - low risk patient	
X142	47.75
X148	61.55

GENERAL PREAMBLE

Code	F
BONE MINERAL DENSITY (BMD) MEASUREMENT (continued)	
Subsequent test - high risk patient	
X149	47.75
X155	61.55
<p>Payment rules:</p> <ol style="list-style-type: none"> 1. Patients are limited to one baseline test (X145 or X146) in their lifetime. 2. Second test – low risk patient (X152/X153) is limited to a maximum of one test rendered not earlier than 36 months following the baseline test (X145/X146). 3. Subsequent test – low risk patient (X142/X148) is <i>not eligible for payment</i> when rendered earlier than 60 months following the second or any subsequent test. 4. Any combination of services described by X152 or X153 that were rendered to a patient between July 1, 2007, and April 1, 2008, for which claims were submitted and paid as insured services under the <i>Health Insurance Act</i> constitutes, a “second test – low risk patient” for the purpose of determining service maximums for a second or subsequent test – low risk patient, and is deemed to have been rendered on July 1, 2010. 5. Any service described by X152 or X153 rendered between April 1, 2008, and July 1, 2010, for which a claim was submitted and paid as an insured service under the <i>Health Insurance Act</i> constitutes a subsequent test – low risk patient for the purpose of determining service maximums for second or subsequent test – low risk patient and is deemed to have been rendered on July 1, 2010. 6. Subsequent test - high risk patients (X149/X155) is limited to a maximum of one test every 12 months unless the ordering physician obtains written prior authorization from a <i>medical consultant</i>. <p>[Commentary: Authorization will be dependent on the referring physician demonstrating that the test is generally accepted as necessary for the patient under the circumstances.]</p> <p>[Commentary:</p> <ol style="list-style-type: none"> 1. Baseline, second test and subsequent tests should be ordered only in accordance with current practice guidelines. In those situations where testing is ordered on a particular patient for reasons that vary from the guidelines, the ordering physician should ensure that the patient’s medical record sufficiently explains the justification for the test in this particular case. 2. In the event a patient with a previous normal baseline test (X145/X146) or second test (X152/X153) or normal subsequent test – low risk patient (X142/X148) meets any of the criteria listed for high risk patients as stated above, the patient would be eligible for subsequent test – high risk patient services (X149/X155) subject to the restriction stated in payment rule #6. 3. The 2002 Clinical Practice Guidelines for the Diagnosis and Management of Osteoporosis in Canada (reviewed in 2006) can be found at http://www.cmaj.ca/cgi/reprint/167/10_suppl/s1.pdf. 4. Individuals under age 65 without one major or two minor risk factors typically do not benefit from BMD measurement.] 	

DIAGNOSTIC RADIOLOGY

Code	F	
MISCELLANEOUS EXAMINATIONS		
X163	Dacrocystogram	33.00
Discogram(s)		
X164	- one or more levels	32.30
X167	Fistula or sinus	23.95
X169	Laminogram, planigram, tomogram	44.50
X170	Laryngogram	32.30
X171	Lymphangiogram	54.65
X192	Mammary ductography	27.95
Mammogram – Signs or Symptoms		
[Commentary: For individuals with identified signs or symptoms or follow-up of established disease.]		
Dedicated equipment		
X184	- unilateral	31.30
X185	- bilateral	41.45
Mammogram – No Signs or Symptoms		
[Commentary: Where the sole reason for the request for a mammogram is for an individual with identified risk factors in accordance with clinical practice guidelines]		
Dedicated equipment		
X172	- unilateral	31.30
X178	- bilateral	41.45
X194	Additional coned views with or without magnification (limit of two per breast) per film	6.65
X201	Breast biopsy specimen x-ray, per specimen	6.65
X150	Mechanical evaluation of knee	28.40
X193	Microradiology of the hands	16.15
X173	Myelogram – spine and/or posterior fossa	38.95
X190	Pantomography	19.80
X154	Penis	17.80
X176	Sialogram	33.25
X177	Skin thickness measurement	17.40
X166	Examination using portable machine “in home” add to first examination only	72.20
<p>Note: X166 does not apply to the use of a portable machine in a hospital. Can only be claimed once per day regardless of the number of people x-rayed in the same “home” including “nursing home”.</p>		

DIAGNOSTIC ULTRASOUND

PREAMBLE

SPECIFIC ELEMENTS

For Facility Fee Component (F)

- A. Preparing the patient for the procedure.
- B. Performing the diagnostic procedure(s).
- C. Making arrangements for any appropriate follow-up care.
- D. Providing records of the results of the procedure to the interpreting physician.
- E. Discussion with, and providing information and advice to, the patient or patient's representative, whether by telephone or otherwise, on matters related to the service.
- F. Preparing and transmitting a written, signed and dated interpretative report of the procedure to the referring physician.
- G. Providing premises, equipment, supplies and personnel for all *specific elements* of the technical components.

OTHER TERMS AND DEFINITIONS

1. Professional and facility fee components are claimed separately. Claims for the facility fee component F are submitted using listed fee code with suffix B. Claims for professional component are submitted using fee code with suffix C (e.g. J102C).
2. A-Mode - implies a one-dimensional ultrasonic measurement procedure.
3. M-Mode - implies a one-dimensional ultrasonic measurement procedure with movement of the trace to record amplitude and velocity of moving echo-producing structures.
4. Scan B-Mode - implies a two-dimensional ultrasonic scanning procedure with a two dimensional display. All ultrasound examinations include a permanent record and interpretative report.
5. All benefits listed apply to unilateral examinations unless otherwise specified. When imaging of only one anatomical area is requested, comparison ultrasound(s) initiated by the interpreting physician or facility are not eligible for payment.
6. Ultrasound of the abdomen, pelvis or breast, rendered in an Independent Health Facility or a hospital in-patient or out-patient department, is insured in accordance with the Health Insurance Act when referred by a registered nurse holding an extended certificate of registration (RN(EC)).
7. Ultrasound for normal, complicated or high risk pregnancy (but not for the postpartum period) rendered in an Independent Health Facility is insured when referred by a midwife who is a member of the College of Midwives of Ontario.

DIAGNOSTIC ULTRASOUND

8. The diagnostic ultrasound benefit includes the generally accepted components of the procedure. For example, where a licensee provides breast ultrasound services, a scan of the axilla is an integral part of the breast imaging exam. The licensee shall not charge any facility fees to the ministry in connection with an additional insured service fee code such as J182 (extremity ultrasound)

9 Where a referring physician requests a single site imaging study (for example, one breast, one limb), any additional imaging of a portion of the anatomy for comparison purposes is not an insured service and shall not be charged to the ministry.

10. Ultrasound of extremity (J182) are to be claimed per limb, not per joint. Scanning two joints on one limb and claiming two services for J182 is incorrect.

11. The practice of routinely submitting claims for more diagnostic ultrasound services than were requested by the referring physician for the majority of patients scanned, will result in a ministry review and potential recovery of funds and/or potential licensing actions. Examples of this unacceptable practice include;

- Bilateral Scans

2 Breasts routinely imaged and billed when only one was requested without the approval of the site radiologist , J127

2, 3, or 4 Extremities routinely imaged and billed when only one or two were requested J182
Axilla scanned and routinely billed as J182 (extremity) during a breast ultrasound [J127 includes scanning of the axilla]

- Routine Addition of scans

Addition of trans vaginal US J138 to a requisition for pelvic US J162

Addition of extremity ultrasound J182 to peripheral vessel assessment, J202

Addition of chest US, J125 to abdominal imaging studies where this is not indicated

Addition of limited pelvis US, J163 to abdominal US, J135, or to limited abdomen, J128

12. Ultrasound services are not insured when rendered in support of in-vitro fertilization services or artificial insemination services.

DIAGNOSTIC ULTRASOUND

Code	F
<u>HEAD AND NECK</u>	
Brain	
J122	48.85
- complete, B-mode	
Echography – ophthalmic (excluding vascular study)	
J102	23.20
- quantitative, A-mode	
J103	45.50
- B-scan immersion	
J107	22.50
- B-scan contact	
J108	23.60
- biometry (Axial length – A-mode)	
Face and/or neck	
J105	48.95
- excluding vascular study	
<p>Note: J105 is <i>not eligible for payment</i> when rendered for ultrasound imaging of the sinus(es).</p>	
<u>THORAX, ABDOMEN AND RETROPERITONEUM</u>	
Thorax	
J125	50.50
Chest masses, pleural effusion – A & B-mode	
Abdomen and Retroperitoneum	
Abdominal scan	
J135	50.50
- complete	
J128	33.25
- limited study (e.g. gallbladder only, aorta only or follow-up study)	
<u>PREGNANCY</u>	
Complete	
J159	50.50
- on or after 16 weeks gestation (maximum one per normal pregnancy)	
J160	50.50
- for high risk pregnancy or complications of pregnancy	
J166	42.90
- multiple gestation, for each additional fetus, to J160	
Gestational age for Maternal Serum Screening Program	
J157	33.25
- before 16 weeks gestation (maximum one per normal pregnancy)	
Limited	
J158	33.25
- for high risk pregnancy or complications of pregnancy	
J167	33.25
- fetal Doppler evaluation of middle cerebral artery and/or ductus venosus, to add J160 or J158,	
<p>Note: J167 is <i>only eligible for payment</i> when rendered by a physician for assessment of fetal anemia or intrauterine growth retardation measuring below the 10th percentile</p>	
J168	40.40
- nuchal translucency for Prenatal Genetic Screening (maximum one per pregnancy)	
J169	34.30
- multiple gestation, for each additional fetus, to J168 add	

DIAGNOSTIC ULTRASOUND

Code	F
<u>PREGNANCY (continued)</u>	
Payment rules: Ultrasound services listed under the headings “Abdomen and Retroperitoneum” or “Pelvis” or “Pregnancy” rendered on the same day to the same patient by any physician as J168 are <i>not eligible for payment</i> .	
<u>PELVIS</u>	
Pelvis	
J162	50.50
- complete*	
J138	50.50
Intracavitary ultrasound* (e.g. transrectal, transvaginal)	
Note: *For ovulation induction purposes, the limit is one per cycle. Additional ultrasounds may be claimed as J164.	
J165	103.50
Transvaginal sonohysterography – may include saline or other intracavitary contrast media except Echovist for demonstration of tubal patency	
J476	241.15
Transvaginal sonohysterography – including Echovist contrast media for demonstration of tubal patency	
Note: J138 and J161 rendered in conjunction with J165 are insured services payable at nil.	
J163	33.25
- limited study – for other than pregnancy	
Intracavitary ultrasound	
J161	33.25
- limited – for other than pregnancy	
J164	25.25
Follicle monitoring studies	
[Commentary: Ultrasound services are not insured when rendered to support in-vitro fertilization services or artificial insemination services.]	
<u>VASCULAR SYSTEM</u>	
Extra-cranial vessel assessment – above the aortic arch	
Bilateral carotid and/or subclavian and/or vertebral arteries only	
J190	44.15
- doppler scan or B scan	
J201	57.00
- duplex scan i.e. simultaneous real time, B-mode imaging and spectral analysis	
Peripheral vessel assessment (distal to inguinal ligament or axilla), artery and/or vein evaluation per extremity. Not to be billed routinely with J190, J191 or J192.	
J193	22.85
- doppler scan or B scan, unilateral	
J202	29.50
- duplex scan i.e. simultaneous real time, B-mode imaging and spectral analysis, unilateral	

DIAGNOSTIC ULTRASOUND

Code		F
VASCULAR SYSTEM (continued)		
Venous assessment		
J198	- bilateral – includes assessment of femoral, popliteal and posterior or tibial veins with appropriate functional manoeuvres and permanent record	7.65
Note: Note to be claimed during surgery or during patient's post-operative stay in hospital.		
Doppler evaluation of organ transplantation		
J205	- arterial and/or venous	22.85
Duplex evaluation of portal hypertension		
J206	- must include doppler interrogation and documentation of superior mesenteric vein, splenic vein, portal veins, hepatic veins and hepatic arteries	22.85
Note: Not to be billed unless study specifically requested by referring physician.		
Duplex assessment of patency obstruction, and flow direction of vascular shunts		
J207	- must include doppler interrogation and documentation of vascular shunts	22.85
Note: Not to be billed unless study specifically requested by referring physician.		
VASCULAR LABORATORY FEES		
Ankle pressure measurements		
J200	- requires a minimum of 4 segmental pressure recordings and/or pulse volume recordings and/or Doppler recordings - unilateral or bilateral	21.10
J196	- with exercise and/or quantitative measurement, to J200	add 8.30
Note: 1. G517 is <i>not eligible for payment</i> in addition to J200. 2. This service is <i>only eligible for payment</i> when the device used produces a hard copy output.		
[Commentary: For ankle pressure determination and ankle-arm index, see G517 under Cardiovascular Diagnostic & Therapeutic Procedures of the Schedule of Benefits.]		
Penile pressure recordings		
J197	- two or more pressures	7.10
Penile Doppler Evaluation		
J199	- Doppler scan	7.10

DIAGNOSTIC ULTRASOUND

Code	F
VASCULAR LABORATORY FEES (continued)	
<p>Note: Penile Doppler is only insured for the following indications:</p> <ol style="list-style-type: none"> 1. priapism; 2. trauma; 3. revascularization; 4. primary erectile dysfunction; or 5. failure of both oral and injectable therapy for erectile dysfunction. <p>[Commentary: Penile Doppler performed for other indications is not an insured service.]</p>	
Transcutaneous tissue	
J203	24.95
J204	13.65
MISCELLANEOUS	
Extremities	
J182	26.40
Breast	
J127	24.55
Scrotal	
J183	48.95
ULTRASONIC GUIDANCE	
<p>SPECIFIC ELEMENTS In addition to the <i>common elements</i>, the components of Ultrasonic Guidance include the following <i>specific elements</i>.</p> <ol style="list-style-type: none"> A. Preparing the patient for the procedure. B. Assisting at the performance of the procedure. C. Making arrangements for follow-up care. D. Discussion with, and providing information and advice to the patient or <i>patient's representative(s)</i>, whether by telephone or otherwise, on matters related to the service. E. Providing premises, equipment, supplies and personnel for all <i>specific elements</i> of the technical and professional components except for the premises for any aspect(s) of A and D of the <i>professional component</i> that is(are) not performed at the place in which the procedure is performed. 	
J149	48.95
<p>Note: J138 and J161 performed during the same visit as J149 is an insured service payable at nil.</p>	

PULMONARY FUNCTION STUDIES

PREAMBLE

SPECIFIC ELEMENTS

For Facility Fee Component (F)

- A. Preparing the patient for the procedure.
- B. Performing the diagnostic procedure
- C. Making arrangements for any appropriate follow-up care.
- D. Providing records of the results of the procedure to the interpreting physician.
- E. Discussion with, and providing information and advice to, the patient or patient's representative, whether by telephone or otherwise, on matters related to the service.
- F. Preparing and transmitting a written, signed and dated interpretive report of the procedure to the referring physician.
- G. Providing premises, equipment, supplies and personnel for all *specific elements* of the technical components.

OTHER TERMS AND DEFINITIONS

1. Professional and facility fee components are claimed separately. Claims for facility fee component F are submitted using listed fee code with suffix B. Claims for professional component P are submitted using listed fee code with suffix C.
2. Each of the following tests designated by an individual code number is considered to be specific and requires individual ordering.
3. Exercise assessment (J315, E450, E451, J316) requires a physician to be in attendance at all times.

PULMONARY FUNCTION STUDIES

Code		F
Functional residual capacity		
J311	- by gas dilution method	16.90
J307	- by body plethysmography	18.10
Note: J311 not to be claimed same patient same day as J307.		
J305	Lung compliance (pressure volume curve of the lung from TLC to FRC)	53.80
J306	Airways resistance by plethysmography or estimated using oesophageal catheter	16.75
J303	Extra pulmonary airways resistance by plethysmography	16.75
J340	Maximum inspiratory and expiratory pressures	2.91
J310	Carbon monoxide diffusing capacity by single breath method	22.15
J308	Carbon dioxide ventilatory response	20.60
Stage I		
J315	Graded exercise to maximum tolerance (exercise must include continuous heart rate, oximetry and ventilation at rest and at each workload)	64.65
E450	- J315 plus J301 or J304 before and/or after exercise	add 13.75
E451	- J315 plus 12 lead E.C.G. done at rest, used for monitoring during the exercise and followed for at least 5 minutes post exercise	add 18.75
Stage II		
J316	Repeated steady state graded exercise (must include heart rate, oximetry, ventilation, VO ₂ , VCO ₂ , BP, ECG, end tidal and mixed Venous CO ₂ at rest, 3 levels of exercise and recovery)	93.20
J330	Assessment of exercise induced asthma (workload sufficient to achieve heart rate 85% of predicted maximum; performance of J301 or J304 before exercise and 5-10 minutes post exercise)	34.55
J319	Blood gas analysis – pH, PO ₂ , PCO ₂ , bicarbonate and base excess	11.65
J318	Arterialized venous blood sample collection (e.g. ear lobe)	3.92
J320	A-a oxygen gradient requiring measurement of RQ by sampling mixed expired gas and using alveolar air equation	28.55
J331	Estimate of shunt (Qs/Qt) breathing pure oxygen	28.55
J313	Mixed venous PCO ₂ , by the rebreathing method	11.65
Oxygen saturation		
J323	- by oximetry at rest, with or without O ₂	4.35
J332	- by oximetry at rest and exercise, or during sleep with or without O ₂	18.20
J334	- J332 with at least two levels of supplemental O ₂	31.65
J336	- with single blind assessment of exercise on room air and with supplemental oxygen	31.65
Note:		
<ol style="list-style-type: none"> 1. J323 is <i>not eligible for payment</i> when rendered with J332, J315, J316 or any overnight sleep study. 2. J332 is <i>not eligible for payment</i> when rendered with J315, J316, or any overnight sleep study. 3. J336 is <i>only eligible for payment</i> for evaluation of a patient to determine eligibility for funding under the Ontario Home Oxygen Program. 4. J336 is not payable in addition to J332 or J334. 		

PULMONARY FUNCTION STUDIES

Code	F
<u>PULMONARY FUNCTION STUDIES (continued)</u>	
Medical record requirements: J323, J332, J334 or J336 are <i>not eligible for payment</i> unless a permanent record of the study is maintained.	
J322	5.50
J333	49.95
J335	53.70
Note: For home/self-care ventilation listing – see Diagnostic and Therapeutic Procedures page J27 of the Schedule of Benefits.	

SLEEP STUDIES

PREAMBLE

SPECIFIC ELEMENTS

For Facility fee Component (F)

- A. Preparing the patient for the procedure.
- B. Performing the diagnostic procedure(s).
- C. Making arrangements for any appropriate follow-up care.
- D. Preparing and providing records of the results of the procedure to the interpreting physician.
- E. Discussion with, and providing information and advice to, the patient or patient's representative, whether by telephone or otherwise, on matters related to the service.
- F. Preparing and transmitting a written, signed and dated interpretative report of the procedure to the referring physician.
- G. Providing premises, equipment, supplies and personnel for all *specific elements* of the technical components.

OTHER TERMS AND DEFINITIONS

SLEEP STUDIES

For the purpose of sleep studies (including overnight sleep studies in non-specialized facilities, overnight sleep studies rendered in specialized facilities and daytime sleep studies),

“CPSO Standards” means the publication of the College of Physicians and Surgeons of Ontario entitled “Independent Health Facilities, Clinical Practice Parameters and Facility Standards, Sleep Medicine” in effect 6 months prior to the date upon which the sleep study was rendered.

“Prior approval” means approved for payment as an insured service, before the service is rendered, by the Ministry of Health following assessment on a case-by-case basis in accordance with all medically relevant criteria.

Sleep studies are subject to limits set out below. Unless otherwise specifically provided, service(s) in excess of these limits are not insured services except when prior approval to exceed the limit is obtained from the Ministry of Health. Despite the foregoing, where prior approval to exceed a limit is not requested from the Ministry of Health but the service would otherwise satisfy one or more of the conditions for which prior approval to exceed the limit is routinely granted (had prior approval been requested) any service in excess of the limit is not eligible for payment.

SLEEP STUDIES

Claims submission instructions:

Submit claims for professional and facility components separately. Submit claims for the facility fee component F using listed fee code with suffix B. Submit claims for professional component using fee code with suffix C (e.g. J890C).

Facility Fee Component

Payment rules:

The facility fee component of the procedure is eligible for payment only if it meets all of the following requirements:

1. A technician is in constant attendance with the patient(s) during the period of the sleep study.
2. The qualifications of technical staff participating in the sleep study comply with the criteria set out in the CPSO Clinical Practice Parameters and Standards.
3. All equipment and test components comply with the criteria set out in the CPSO Standards.

Medical record requirements:

Records of the facility fee component must conform to the standards for facilities and facility operators (including records required prior to data analysis) as set out in the CPSO Clinical Practice Parameters and Standards, or the facility fee component is not eligible for payment.

SLEEP STUDIES

Code	F
<p>OVERNIGHT SLEEP STUDIES</p> <p>For the purpose of sleep studies (including overnight sleep studies and daytime sleep studies),</p> <p>“CPSO Standards” means the publication of the College of Physicians and Surgeons of Ontario entitled “Independent Health Facilities, Clinical Practice Parameters and Facility Standards, Sleep Medicine” in effect 6 months prior to the date upon which the sleep study was rendered.</p> <p>“prior approval” means approved for payment as an insured service, before the service is rendered, by the Ministry of Health following assessment on a case-by-case basis in accordance with all medically relevant criteria.</p> <p>Terms and Conditions</p> <p>Facility fees for sleep studies meeting the eligibility parameters are payable under the Independent Health Facilities Act and are listed in the Schedule of Facility Fees.</p> <p>Sleep studies are subject to limits set out below. Unless otherwise specifically provided, service(s) in excess of these limits are not insured services except when prior approval to exceed the limit is obtained from the Ministry of Health. Despite the foregoing, where prior approval to exceed a limit is not requested from the Ministry of Health but the service would otherwise satisfy one or more of the conditions for which prior approval to exceed the limit is routinely granted (had prior approval been requested) any service in excess of the limit is not eligible for payment.</p> <p>[Commentary: Services rendered in excess of a maximum are not eligible for payment.]</p> <p>IHF Facility Fee Payment rules:</p> <p>The facility fee for the procedure is eligible for payment only if it meets all of the following requirements:</p> <ol style="list-style-type: none"> 1. It satisfies the conditions set out under “Sleep Studies Services Rendered at a licensed Independent Health Facility (IHF)”. 2. It is rendered at a licensed IHF. 3. A technician is in constant attendance with the patient(s) during the period of the sleep study. 4. The qualifications of technical staff participating in the sleep study comply with the criteria set out in the CPSO Standards. 5. All equipment and test components comply with the criteria set out in the CPSO Standards. 	

SLEEP STUDIES

Code		F
	<p>“Sleep Studies Services Rendered at a licensed Independent Health Facility (IHF)”.</p> <p>A. Incomplete Overnight Sleep Studies</p> <p>If the recording does not contain information sufficient for a diagnostic interpretation as determined in accordance with generally accepted standards as set out in the CPSO Standards, the professional fee is <i>not eligible for payment</i> and the service constitutes one of the following, as determined by time in bed (total study time):</p>	
J898	Sleep study less than 1 hour	95.95
J899	Sleep study between 1 and 4 hours	191.95
J990	Sleep study more than 4 hours	383.85
	<p>Payment rules:</p> <ol style="list-style-type: none"> 1. A maximum of one of any of J898, J899 and J990 is eligible for payment, per patient, per facility, per <i>12 month period</i>. 2. J898, J899 and J990 are not included in the limits for overnight studies set out below. <p>B. Overnight Sleep Studies in Independent Health Facilities</p> <p>Level 1</p> <p>Is a overnight sleep study with continuous monitoring of oxygen saturation, ECG and Ventilation (airflow and respiratory effort) and additional monitoring to stage sleep (including all of the following: EEG, EOG and sub-mental EMG).</p> <p style="text-align: center;">Initial Diagnostic Study</p> <p>“Initial Diagnostic Study” means the first overnight sleep study rendered to an insured person as an insured service in Ontario for the purpose of establishing the diagnosis of a sleep disorder (and includes a split night study). Every overnight diagnostic sleep study rendered before July 1, 2010, for which a claim was submitted and paid as an insured service under the <i>Health Insurance Act</i> constitutes an “initial diagnostic study” and is deemed to have been rendered on July 1, 2010.</p>	
	Initial Diagnostic Study – Level 1	
J896	- diagnostic study	383.85
	<p>Note:</p> <ol style="list-style-type: none"> 1. A maximum of one initial diagnostic study is eligible for payment per patient per lifetime. 2. All subsequent overnight sleep studies constitute “repeat diagnostic” or “therapeutic” studies. 	

SLEEP STUDIES

Code	F
<p style="text-align: center;">Repeat Diagnostic Study</p> <p>“Repeat Diagnostic Study” means an overnight diagnostic sleep study rendered:</p> <ul style="list-style-type: none"> a. for the purpose of obtaining a second opinion at a different facility than the facility where the preceding study was rendered, provided that the following conditions are met: <ul style="list-style-type: none"> i. prior to the repeat diagnostic study, the patient has been assessed by a physician who practices sleep medicine at the different facility, <p>[Commentary: The different facility requirement above applies to a repeat diagnostic study rendered at a hospital, a hospital off-site premise or an independent health facility.]</p> <ul style="list-style-type: none"> ii. where the previous study was rendered at an independent health facility and the repeat diagnostic study is rendered at a different independent health facility (the “different facility”) than the independent health facility where the preceding study was rendered (the “first facility”), neither the owner nor the operator of the different facility is, at the time the repeat study is rendered, an associate of the owner or operator of the first facility, where “associate” has the same meaning as in the <i>Independent Health Facilities Act</i>; <p>OR</p> <ul style="list-style-type: none"> b. for one or more of the following purposes, after pre-study assessment by a physician practicing sleep medicine: <ul style="list-style-type: none"> i. re-evaluation of a previous negative or inconclusive diagnostic sleep study as indicated by persistent or progressive symptoms; ii. re-evaluation, other than primarily for Positive Airway Pressure therapy (PAP) adjustment, of patients previously diagnosed with a primary sleep disorder in which there has been symptom development suggesting another co-morbid sleep disorder; or iii. re-evaluation of patients with an established diagnosis of a sleep disorder other than a sleep related breathing disorder who have significant symptom progression or non-response to therapy <p>[Commentary: 1. In the case of patients with previously diagnosed sleep related breathing disorders, although PAP treatment may be adjusted during a repeat study, a repeat study is <i>not eligible for payment</i> if rendered primarily for PAP treatment adjustment. 2. Examples of sleep disorders other than a sleep related breathing disorder are Narcolepsy, Idiopathic hypersomnia and Periodic Limb Movement Disorder.]</p>	
J897 - diagnostic study	383.85

SLEEP STUDIES

Code	F
Repeat Diagnostic Study – Level 1 (continued)	
<p>Payment rules:</p> <ol style="list-style-type: none"> 1. Repeat diagnostic studies are limited to one per patient, per facility, per <i>12 month period</i> except where prior approval has been given. 2. Repeat diagnostic studies performed in the same facility that performed the initial diagnostic study are <i>not eligible for payment</i> in the <i>12 month period</i> following an initial diagnostic study except where prior approval has been given. <p>Therapeutic study</p> <p>“Therapeutic Study” means a sleep study rendered after pre-study assessment by a physician practicing sleep medicine, for any of the following purposes:</p> <ol style="list-style-type: none"> a. To establish optimal settings for nasal positive airway pressure therapy (CPAP/BiPAP/ASV etc.) and/or oxygen therapy for sleep related breathing disorders; <p>[Commentary: Examples of sleep related breathing disorders are obstructive sleep apnea syndrome (OSAS), central sleep apnea syndrome (CSAS), Cheyne-Stokes breathing, complex sleep apnea syndrome, or hypoventilation syndromes.]</p> <ol style="list-style-type: none"> b. To evaluate the response to surgical procedures for the treatment of OSAS; c. To determine the efficacy of oral appliance therapy for OSAS; d. To evaluate the efficacy of positional therapy for the treatment of OSAS; e. To evaluate the efficacy of substantial weight loss for the treatment of OSAS; or f. To titrate ventilatory settings for patients with respiratory control disorders, neuromuscular or neurodegenerative diseases. 	
Therapeutic Study for Sleep Related Breathing Disorders – Level 1	
J895	383.85
<p>- therapeutic study</p>	
<p>Payment rules:</p> <ol style="list-style-type: none"> 1. There is a limit of one therapeutic study (J895) per patient during any two consecutive <i>12 month</i> periods except where prior approval has been given. 2. J895 rendered to the same patient during the same 12 hour period as J896 or J897 is <i>not eligible for payment</i>. 	
<p>Note:</p> <ol style="list-style-type: none"> 1. For payment purposes, repeat diagnostic studies or therapeutic studies for indications or in circumstances other than listed above, or in excess of the limits set out below, require prior approval. 2. A repeat diagnostic study rendered without the required pre-study assessment by a physician practicing sleep medicine, is <i>not eligible for payment</i>. 	

SLEEP STUDIES

Code	F	
<u>Therapeutic Study for Sleep Related Breathing Disorders – Level 1 (continued)</u>		
<p>3. A therapeutic study rendered without a pre-study assessment by a physician practicing sleep medicine is <i>not eligible for payment</i> except:</p> <ul style="list-style-type: none"> a. For the therapeutic study that immediately follows an initial diagnostic or repeat diagnostic study where: <ul style="list-style-type: none"> i. the time interval is such that it is unlikely the clinical circumstances of the patient has changed; and ii. the physician practicing sleep medicine has previously assessed the patient and documented the applicable decisions with respect to the performance of the therapeutic study; or b. In exceptional circumstances where the physician can demonstrate to the ministry upon request that the CPSO standards are satisfied with the use of a clinical protocol or approved medical directive. 		
<p>[Commentary:</p> <ol style="list-style-type: none"> 1. An example of an exceptional circumstance may be where a patient is required to travel a long distance to a sleep facility and requires an initial diagnostic or repeat diagnostic study followed by a therapeutic study on a subsequent night. For payment purposes, a pre-study assessment by a physician practicing sleep medicine is not required provided the therapeutic study is rendered in accordance with a clinical protocol or medical directive that has been approved by an authority other than a physician affiliated with the sleep facility (e.g. a Medical Advisory Committee for a sleep clinic affiliated with a hospital). The physician should be prepared to provide any necessary supporting documentation to the ministry upon request. 2. Prior approval, where required, will typically be dependent on the physician demonstrating that the study is generally accepted as necessary for the patient under the circumstances. 3. Sleep studies that require prior approval also require a pre-study assessment by a physician practicing sleep medicine. It is this assessment upon which the request for prior approval is considered. 4. Prior approval requires a written request accompanied by supporting documentation including the pre-study assessment and the relevant previous sleep study reports. 5. Split-night sleep studies are claimed as J896 or J897 only, as appropriate to the study rendered.] 		
C. Daytime Sleep Studies		
J893	Multiple sleep latency test	71.40
J894	Maintenance of wakefulness test	71.40
<p>Payment rules:</p> <ol style="list-style-type: none"> 1. J894 rendered to same patient same <i>day</i> as J893 is <i>not eligible for payment</i>. 2. A maximum of one J893 and a maximum of one J894 are payable per <i>12 month period</i> per facility per patient. 		

SLEEP STUDIES

Code	F
<u>Daytime sleep studies (continued)</u>	
3. If the recording does not contain information sufficient for a diagnostic interpretation as determined in accordance with CPSO standards, the service is <i>not eligible for payment.</i>	