

Medical Policy



Title: Intensity Modulated Radiation Therapy (IMRT)

See also: *Stereotactic Radiosurgery and Radiotherapy medical policy*

PRE-DETERMINATION of services for FEP is required effective January 1, 2010.

http://www.bcbsks.com/CustomService/Forms/pdf/15-17_predeterm_request_frm.pdf

Professional

Original Effective Date: January 1, 2002
Revision Date(s): January 30, 2009;
January 1, 2010; October 11, 2011;
July 13, 2012; October 15, 2012;
December 11, 2013
Current Effective Date: October 15, 2012

Institutional

Original Effective Date: March 2, 2009
Revision Date(s): January 1, 2010;
October 11, 2011; July 13, 2012;
October 15, 2012; December 11, 2013
Current Effective Date: October 15, 2012

State and Federal mandates and health plan member contract language, including specific provisions/exclusions, take precedence over Medical Policy and must be considered first in determining eligibility for coverage. To verify a member's benefits, contact [Blue Cross and Blue Shield of Kansas Customer Service](#).

The BCBSKS Medical Policies contained herein are for informational purposes and apply only to members who have health insurance through BCBSKS or who are covered by a self-insured group plan administered by BCBSKS. Medical Policy for FEP members is subject to FEP medical policy which may differ from BCBSKS Medical Policy.

The medical policies do not constitute medical advice or medical care. Treating health care providers are independent contractors and are neither employees nor agents of Blue Cross and Blue Shield of Kansas and are solely responsible for diagnosis, treatment and medical advice.

If your patient is covered under a different Blue Cross and Blue Shield plan, please refer to the Medical Policies of that plan.

DESCRIPTION

Intensity modulated radiation therapy (IMRT) has been proposed as a method of radiation therapy that allows adequate radiation therapy to the tumor while minimizing the radiation dose to surrounding normal tissues and critical structures.

Radiation Techniques

Conventional external-beam radiation therapy.

Over the past several decades, methods to plan and deliver radiation therapy have evolved in ways that permit more precise targeting of tumors with complex geometries. Most early trials used 2-dimensional treatment planning, based on flat images and radiation beams with cross-sections of uniform intensity that were sequentially aimed at the tumor along 2 or 3 intersecting axes. Collectively, these methods are termed “conventional external beam radiation therapy”.

3-dimensional conformal radiation (3D-CRT).

Treatment planning evolved by using 3-dimensional images, usually from computed tomography (CT) scans, to delineate the boundaries of the tumor and discriminate tumor tissue from adjacent normal tissue and nearby organs at risk for radiation damage. Computer algorithms were developed to estimate cumulative radiation dose delivered to each volume of interest by summing the contribution from each shaped beam. Methods also were developed to position the patient and the radiation portal reproducibly for each fraction and immobilize the patient, thus maintaining consistent beam axes across treatment sessions. Collectively, these methods are termed 3-dimensional conformal radiation therapy (3D-CRT).

Intensity-modulated radiation therapy (IMRT).

IMRT, which uses computer software, CT images, and magnetic resonance imaging (MRI), offers better conformality than 3D-CRT as it is able to modulate the intensity of the overlapping radiation beams projected on the target and to use multiple-shaped treatment fields. It uses a device (a multileaf collimator, MLC) which, coupled to a computer algorithm, allows for “inverse” treatment planning. The radiation oncologist delineates the target on each slice of a CT scan and specifies the target’s prescribed radiation dose, acceptable limits of dose heterogeneity within the target volume, adjacent normal tissue volumes to avoid, and acceptable dose limits within the normal tissues. Based on these parameters and a digitally reconstructed radiographic image of the tumor and surrounding tissues and organs at risk, computer software optimizes the location, shape, and intensities of the beams ports, to achieve the treatment plan’s goals.

Increased conformality may permit escalated tumor doses without increasing normal tissue toxicity and thus may improve local tumor control, with decreased exposure to surrounding normal tissues, potentially reducing acute and late radiation toxicities. Better dose homogeneity within the target may also improve local tumor control by avoiding underdosing within the tumor and may decrease toxicity by avoiding overdosing.

Since most tumors move as patients breathe, dosimetry with stationary targets may not accurately reflect doses delivered within target volumes and adjacent tissues in patients. Furthermore, treatment planning and delivery are more complex, time-consuming, and labor-intensive for IMRT than for 3D-CRT. Thus, clinical studies must test whether IMRT improves tumor control or reduces acute and late toxicities when compared with 3D-CRT.

Methodologic issues with IMRT studies

Multiple-dose planning studies have generated 3D-CRT and IMRT treatment plans from the same scans, then compared predicted dose distributions within the target and in adjacent organs at risk. Results of such planning studies show that IMRT improves on 3D-CRT with respect to conformality to, and dose homogeneity within, the target. Dosimetry using stationary targets generally confirms these predictions. Thus, radiation oncologists hypothesized that IMRT may improve treatment outcomes compared with those of 3D-CRT. However, these types of studies offer indirect evidence

on treatment benefit from IMRT, and it is difficult to relate results of dosing studies to actual effects on health outcomes.

Comparative studies of radiation-induced side effects from IMRT versus alternative radiation delivery are probably the most important type of evidence in establishing the benefit of IMRT. Such studies would answer the question of whether the theoretical benefit of IMRT in sparing normal tissue translates into real health outcomes. Single-arm series of IMRT can give some insights into the potential for benefit, particularly if an adverse effect that is expected to occur at high rates is shown to decrease by a large amount. Studies of treatment benefit are also important to establish that IMRT is at least as good as other types of delivery, but in the absence of such comparative trials, it is likely that benefit from IMRT is at least as good as with other types of delivery.

POLICY

IMRT is considered **medically necessary** for the following indications:

- A. Prostate cancer for dose escalation >75 Gy of the prostate and for post operative radiation of the prostate to a dose of at least 6300 cGy.
- B. Head and neck cancer, with the exception of patients with early stage larynx cancer (stage I and II).
- C. Cancer involving the CNS.
- D. Carcinoma of the anus or vulva.
- E. Anaplastic thyroid cancer.
- F. Carcinoma of the cervix
- G. Whole pelvic radiotherapy for gynecologic malignancies.
- H. Pediatric tumors (e.g., Ewing Sarcoma, Wilms' Tumor).
- I. Breast cancer, when at least one of the following is met:
 1. Heart—3D results in $\geq 25\%$ of heart receiving $\geq 30\text{Gy}$;
OR
 2. Lung—3D results in $\geq 30\%$ of ipsilateral lung receiving $\geq 20\text{ Gy}$,
OR
3D results in $\geq 20\%$ of combined lung volume receiving $\geq 20\text{Gy}$;
OR
 3. Skin / Soft Tissue—3D results in $\geq 5\%$ of intended breast receiving $\geq 7\%$ of prescribed dose
OR
Medial lesion where 3D results in $\geq 10\%$ of contralateral breast receiving $\geq 10\text{Gy}$.

- J. Esophagus, Stomach, Pancreas, Hepatobiliary Tract, Rectum, Colon, Small Bowel, when at least one of the following is met:
1. Heart—3D result in $\geq 50\%$ of heart receiving ≥ 30 Gy,
OR
 2. Lung—3D results in $\geq 30\%$ of combined lung volume receiving $\geq 20\%$ Gy,
OR
Mean lung dose ≥ 20 Gy;
OR
 3. Spinal Cord—3D result in any portion of the spinal cord receiving a dose above 45 Gy;
OR
 4. Liver—3D results in $\geq 60\%$ of liver volume receiving ≥ 30 Gy,
OR
Mean liver dose ≥ 32 Gy;
OR
 5. Kidney—3D results in $\geq 33\%$ of combined kidney volume receiving ≥ 20 Gy (two functional kidneys are present);
OR
For one functioning kidney or kidney transplant, IMRT provides a lower dose than achievable with 3D;
OR
 6. Small Intestine—3D results in ≥ 195 cc of small intestine receiving ≥ 45 Gy;
OR
 7. Stomach—3D results in $\geq 10\%$ of stomach receiving ≥ 45 Gy
OR
5% receiving ≥ 50 Gy;
OR
 8. Femoral head—3D results in a femoral head receiving ≥ 45 Gy.
- K. Lung, when at least one of the following is met:
1. Heart—3D results in $\geq 50\%$ of heart receiving ≥ 30 Gy;
OR
 2. Lung—3D result in $\geq 30\%$ of non-cancerous combined lung volume receiving ≥ 20 Gy
- L. Lymphomas or Sarcomas of Retroperitoneum, Chest Wall and Thorax, when at least one of the following is met:
1. Heart—3D results in $\geq 50\%$ of heart receiving ≥ 30 Gy;
OR
 2. Lung—3D results in $\geq 30\%$ of combined lung volume receiving ≥ 20 Gy
OR

- Mean lung dose of ≥ 20 Gy;
OR
3. Spinal cord—3D results in any portion of the spinal cord receiving a dose above 45 Gy
OR
 4. Liver—3D results in 60% of liver volume receiving ≥ 30 Gy
OR
 Mean liver dose ≥ 32 Gy;
OR
 5. Femoral head—3D results in a femoral head receiving ≥ 45 Gy;
OR
 6. Small intestine—3D results in ≥ 195 cc of small intestine receiving ≥ 45 Gy;
OR
 7. Stomach—3D results in $\geq 10\%$ of stomach receiving ≥ 45 Gy
OR
 5% receiving ≥ 50 Gy;
OR
 8. Rectosigmoid—3D results in $\geq 60\%$ of rectosigmoid area receiving ≥ 30 Gy;
OR
 9. Bladder—3D results in $\geq 35\%$ of bladder receiving ≥ 45 Gy;
OR
 10. Kidney—3D results in 33% of combined kidney volume receiving ≥ 20 Gy (two functional kidneys are present)
OR
 For one functioning kidney or kidney transplant IMRT provides a lower dose than achievable with 3D.
- M. Sarcomas of the Extremities, when at least one of the following is met:
1. Head / Neck—IMRT covered if head and neck structures would receive any radiation via 3D;
OR
 2. Femur—3D results in $\geq 50\%$ of contiguous femur cortex receiving ≥ 50 Gy
- N. Individuals who require repeat irradiation of a field that has received prior irradiation.
- O. Radiosensitive tumors where critical structures cannot be adequately protected with standard 3D conformal radiotherapy. Medical necessity for the use of IMRT for these other indications will be considered individually and will require supporting records from the treating radiation oncologist including the 3-D dose volume histogram documenting the need for IMRT.

The American Society for Therapeutic Radiology and Oncology (ASTRO) has a model policy which describes the indications for IMRT:

"IMRT is not a replacement therapy for conventional or three-dimensional conformal radiation therapy methods. IMRT is considered reasonable and necessary in instances where sparing the surrounding normal tissue is of added benefit and at least one of the following conditions is met:

1. The target volume is in close proximity to critical structures that must be protected.
2. The volume of interest must be covered with narrow margins to adequately protect immediately adjacent structures.
3. An immediately adjacent area has been previously irradiated and abutting portals must be established with high precision.
4. The target volume is concave or convex, and critical normal tissues are within or around that convexity or concavity.
5. Dose escalation is planned to deliver radiation doses in excess of those commonly utilized for similar tumor with conventional treatments."

P. Other applications of IMRT are considered **not medically necessary**.

DOCUMENTATION

As recommended by ASTRO, the IMRT treatment record must include:

1. The reasonable and necessary requirements as outlined in the Policy section.
2. The prescription defining the dose to the target and the dose constraints to the nearby critical structures.
3. A note of medical necessity for IMRT, by the treating physician.
4. Signed IMRT inverse plan that meets prescribed dose constraints for the planning target volume (PTV) and surrounding normal tissue.
5. The target verification methodology including the following:
 - a. Documentation of the clinical treatment volume (CTV) and the planning target volume (PTV).
 - b. Documentation of immobilization and patient positioning.
6. Independent basic dose calculations of monitor units performed for each beam before the patient's first treatment.
7. Documentation of fluence distributions (re-computed and measured in a phantom or dosimetry measuring device).
8. Identification of structures that transverse high-and low-dose regions created by respiration. Voluntary breath-holding alone is not a satisfactory solution for accounting for organ motion.

CODING

The following codes for treatment and procedures applicable to this policy are included below for informational purposes. Inclusion or exclusion of a procedure, diagnosis or device code(s) does not constitute or imply member coverage or provider reimbursement. Please refer to the member's contract benefits in effect at the time of service to determine coverage or non-coverage of these services as it applies to an individual member.

CPT/HCPCS

- 77300 Basic radiation dosimetry calculation, central axis depth dose calculation, TDF, NSD, gap calculation, off axis factor, tissue inhomogeneity factors, calculation of non- ionizing radiation surface and depth dose, as required during course of treatment, only when prescribed by the treating physician
- 77301 Intensity modulated radiotherapy plan, including dose-volume histograms for target and critical structure partial tolerance specifications
- 77332 Treatment devices, design and construction; simple (simple block, simple bolus)
- 77333 Treatment devices, design and construction; intermediate (multiple blocks, stents, bite blocks, special bolus)
- 77334 Treatment devices, design and construction; complex (irregular blocks, special shields, compensators, wedges, molds or casts)
- 77338 Multi-leaf collimator (MLC) device(s) for intensity modulated radiation therapy (IMRT), design and construction per IMRT plan
- 77418 Intensity modulated treatment delivery, single or multiple fields/arcs, via narrow spatially and temporally modulated beams, binary, dynamic MLC, per treatment session
- 0073T Compensator-based beam modulation treatment delivery of inverse planned treatment using three or more high resolution (milled or cast) compensator convergent beam modulated fields, per treatment session

DIAGNOSIS

- 140.0- Malignant neoplasm of lip, oral cavity, and pharynx
- 149.9
- 154.2- Malignant neoplasm of anus
- 154.3
- 160.0- Malignant neoplasm of nasal cavities, middle ear, and accessory sinuses
- 160.9
- 161.0- Malignant neoplasm of larynx
- 161.9
- 170.0- Malignant neoplasm of bones of skull and face, mandible
- 170.1
- 171.0 Malignant neoplasm of connective and other soft tissue of head, face, and neck
- 172.0- Malignant melanoma of lip, eyelid, ear and external auditory canal, other and
- 172.4 unspecified parts of face, scalp and neck
- 173.0- Other malignant neoplasm of skin of lip, eyelid, ear and external auditory canal, other and
- 173.4 unspecified parts of face, scalp and neck
- 174.0- Malignant neoplasm of female breast
- 174.9
- 180.-0 Malignant neoplasm of cervix uteri, code range
- 180.9

182.0-	Malignant neoplasm of body of uterus, code range
182.8	
185	Malignant neoplasm of prostate
190.0-	Malignant neoplasm of eye
190.9	
191.0-	Malignant neoplasm of brain
191.9	
192.0-	Malignant neoplasm of other and unspecified parts of nervous system
192.9	
193	Malignant neoplasm of thyroid gland
194.1-	Malignant neoplasm of parathyroid, pituitary, pineal gland, carotid body
194.5	
195.0	Malignant neoplasm of other and ill-defined sites-head, face, neck
196.0	Malignant neoplasm of lymph nodes of head, face and neck
198.3	Secondary malignant neoplasm of brain and spinal cord
200.01	Reticulosarcoma involving lymph nodes of head, face, and neck
200.11	Lymphosarcoma involving lymph nodes of head, face, and neck
225.0-	Benign neoplasm of brain and other parts of nervous system
225.9	
227.1-	Benign neoplasm of parathyroid gland, pituitary gland and craniopharyngeal duct
227.6	(pouch), pineal gland, carotid body, aortic body and other paraganglia
236.5	Neoplasm of uncertain behavior of genitourinary organs-prostate
237.0	Neoplasm of uncertain behavior of pituitary gland and craniopharyngeal duct
237.1	Neoplasm of uncertain behavior of pineal gland
237.5	Neoplasm of uncertain behavior of endocrine glands and nervous system-brain and spinal cord
237.6	Neoplasm of uncertain behavior of meninges
V10.40	Personal history of malignant neoplasm of genital organs; female genital organ. unspecified
V10.41	Personal history of malignant neoplasm of genital organs; cervix uteri

ICD-10 Diagnosis (*Effective October 1, 2014*)

C00.0	Malignant neoplasm of external upper lip
C00.1	Malignant neoplasm of external lower lip
C00.3	Malignant neoplasm of upper lip, inner aspect
C00.4	Malignant neoplasm of lower lip, inner aspect
C00.8	Malignant neoplasm of overlapping sites of lip
C01	Malignant neoplasm of base of tongue
C02.0	Malignant neoplasm of dorsal surface of tongue
C02.1	Malignant neoplasm of border of tongue
C02.2	Malignant neoplasm of ventral surface of tongue
C02.3	Malignant neoplasm of anterior two-thirds of tongue, part unspecified
C02.4	Malignant neoplasm of lingual tonsil
C02.8	Malignant neoplasm of overlapping sites of tongue
C03.0	Malignant neoplasm of upper gum
C03.1	Malignant neoplasm of lower gum
C04.0	Malignant neoplasm of anterior floor of mouth
C04.1	Malignant neoplasm of lateral floor of mouth
C04.8	Malignant neoplasm of overlapping sites of floor of mouth
C05.0	Malignant neoplasm of hard palate

- C05.1 Malignant neoplasm of soft palate
- C05.2 Malignant neoplasm of uvula
- C05.8 Malignant neoplasm of overlapping sites of palate
- C06.0 Malignant neoplasm of cheek mucosa
- C06.1 Malignant neoplasm of vestibule of mouth
- C06.2 Malignant neoplasm of retromolar area
- C06.89 Malignant neoplasm of overlapping sites of other parts of mouth
- C07 Malignant neoplasm of parotid gland
- C08.0 Malignant neoplasm of submandibular gland
- C08.1 Malignant neoplasm of sublingual gland
- C08.9 Malignant neoplasm of major salivary gland, unspecified
- C09.0 Malignant neoplasm of tonsillar fossa
- C09.1 Malignant neoplasm of tonsillar pillar (anterior) (posterior)
- C09.8 Malignant neoplasm of overlapping sites of tonsil
- C10.0 Malignant neoplasm of vallecula
- C10.1 Malignant neoplasm of anterior surface of epiglottis
- C10.2 Malignant neoplasm of lateral wall of oropharynx
- C10.3 Malignant neoplasm of posterior wall of oropharynx
- C10.4 Malignant neoplasm of branchial cleft
- C10.8 Malignant neoplasm of overlapping sites of oropharynx
- C11.0 Malignant neoplasm of superior wall of nasopharynx
- C11.1 Malignant neoplasm of posterior wall of nasopharynx
- C11.2 Malignant neoplasm of lateral wall of nasopharynx
- C11.3 Malignant neoplasm of anterior wall of nasopharynx
- C11.8 Malignant neoplasm of overlapping sites of nasopharynx
- C12 Malignant neoplasm of pyriform sinus
- C13.0 Malignant neoplasm of postcricoid region
- C13.1 Malignant neoplasm of aryepiglottic fold, hypopharyngeal aspect
- C13.2 Malignant neoplasm of posterior wall of hypopharynx
- C13.8 Malignant neoplasm of overlapping sites of hypopharynx
- C14.0 Malignant neoplasm of pharynx, unspecified
- C14.2 Malignant neoplasm of Waldeyer's ring
- C14.8 Malignant neoplasm of overlapping sites of lip, oral cavity and pharynx
- C21.0 Malignant neoplasm of anus, unspecified
- C21.1 Malignant neoplasm of anal canal
- C30.0 Malignant neoplasm of nasal cavity
- C30.1 Malignant neoplasm of middle ear
- C31.0 Malignant neoplasm of maxillary sinus
- C31.1 Malignant neoplasm of ethmoidal sinus
- C31.2 Malignant neoplasm of frontal sinus
- C31.3 Malignant neoplasm of sphenoid sinus
- C31.8 Malignant neoplasm of overlapping sites of accessory sinuses
- C32.0 Malignant neoplasm of glottis
- C32.1 Malignant neoplasm of supraglottis
- C32.2 Malignant neoplasm of subglottis
- C32.3 Malignant neoplasm of laryngeal cartilage
- C32.8 Malignant neoplasm of overlapping sites of larynx
- C41.0 Malignant neoplasm of bones of skull and face
- C41.1 Malignant neoplasm of mandible

- C43.0 Malignant melanoma of lip
- C43.11 Malignant melanoma of right eyelid, including canthus
- C43.12 Malignant melanoma of left eyelid, including canthus
- C43.21 Malignant melanoma of right ear and external auricular canal
- C43.22 Malignant melanoma of left ear and external auricular canal
- C43.31 Malignant melanoma of nose
- C43.39 Malignant melanoma of other parts of face
- C43.4 Malignant melanoma of scalp and neck
- C47.0 Malignant neoplasm of peripheral nerves of head, face and neck
- C49.0 Malignant neoplasm of connective and soft tissue of head, face and neck
- D03.0 Melanoma in situ of lip
- D03.11 Melanoma in situ of right eyelid, including canthus
- D03.12 Melanoma in situ of left eyelid, including canthus
- D03.21 Melanoma in situ of right ear and external auricular canal
- D03.22 Melanoma in situ of left ear and external auricular canal
- D03.39 Melanoma in situ of other parts of face
- D03.4 Melanoma in situ of scalp and neck
- C44.00 Unspecified malignant neoplasm of skin of lip
- C44.01 Basal cell carcinoma of skin of lip
- C44.02 Squamous cell carcinoma of skin of lip
- C44.09 Other specified malignant neoplasm of skin of lip
- C44.102 Unspecified malignant neoplasm of skin of right eyelid, including canthus
- C44.109 Unspecified malignant neoplasm of skin of left eyelid, including canthus
- C44.112 Basal cell carcinoma of skin of right eyelid, including canthus
- C44.119 Basal cell carcinoma of skin of left eyelid, including canthus
- C44.122 Squamous cell carcinoma of skin of right eyelid, including canthus
- C44.129 Squamous cell carcinoma of skin of left eyelid, including canthus
- C44.191 Other specified malignant neoplasm of skin of unspecified eyelid, including canthus
- C44.192 Other specified malignant neoplasm of skin of right eyelid, including canthus
- C44.199 Other specified malignant neoplasm of skin of left eyelid, including canthus
- C44.202 Unspecified malignant neoplasm of skin of right ear and external auricular canal
- C44.209 Unspecified malignant neoplasm of skin of left ear and external auricular canal
- C44.212 Basal cell carcinoma of skin of right ear and external auricular canal
- C44.219 Basal cell carcinoma of skin of left ear and external auricular canal
- C44.222 Squamous cell carcinoma of skin of right ear and external auricular canal
- C44.229 Squamous cell carcinoma of skin of left ear and external auricular canal
- C44.292 Other specified malignant neoplasm of skin of right ear and external auricular canal
- C44.299 Other specified malignant neoplasm of skin of left ear and external auricular canal
- C44.301 Unspecified malignant neoplasm of skin of nose
- C44.309 Unspecified malignant neoplasm of skin of other parts of face
- C44.311 Basal cell carcinoma of skin of nose
- C44.319 Basal cell carcinoma of skin of other parts of face
- C44.321 Squamous cell carcinoma of skin of nose
- C44.329 Squamous cell carcinoma of skin of other parts of face
- C44.391 Other specified malignant neoplasm of skin of nose
- C44.399 Other specified malignant neoplasm of skin of other parts of face
- C44.40 Unspecified malignant neoplasm of skin of scalp and neck
- C44.41 Basal cell carcinoma of skin of scalp and neck
- C44.42 Squamous cell carcinoma of skin of scalp and neck

- C44.49 Other specified malignant neoplasm of skin of scalp and neck
- C61 Malignant neoplasm of prostate
- C69.01 Malignant neoplasm of right conjunctiva
- C69.02 Malignant neoplasm of left conjunctiva
- C69.11 Malignant neoplasm of right cornea
- C69.12 Malignant neoplasm of left cornea
- C69.21 Malignant neoplasm of right retina
- C69.22 Malignant neoplasm of left retina
- C69.31 Malignant neoplasm of right choroid
- C69.32 Malignant neoplasm of left choroid
- C69.41 Malignant neoplasm of right ciliary body
- C69.42 Malignant neoplasm of left ciliary body
- C69.51 Malignant neoplasm of right lacrimal gland and duct
- C69.52 Malignant neoplasm of left lacrimal gland and duct
- C69.61 Malignant neoplasm of right orbit
- C69.62 Malignant neoplasm of left orbit
- C69.81 Malignant neoplasm of overlapping sites of right eye and adnexa
- C69.82 Malignant neoplasm of overlapping sites of left eye and adnexa
- C69.91 Malignant neoplasm of unspecified site of right eye
- C69.92 Malignant neoplasm of unspecified site of left eye
- C70.0 Malignant neoplasm of cerebral meninges
- C70.1 Malignant neoplasm of spinal meninges
- C70.9 Malignant neoplasm of meninges, unspecified
- C71.0 Malignant neoplasm of cerebrum, except lobes and ventricles
- C71.1 Malignant neoplasm of frontal lobe
- C71.2 Malignant neoplasm of temporal lobe
- C71.3 Malignant neoplasm of parietal lobe
- C71.4 Malignant neoplasm of occipital lobe
- C71.5 Malignant neoplasm of cerebral ventricle
- C71.6 Malignant neoplasm of cerebellum
- C71.7 Malignant neoplasm of brain stem
- C71.8 Malignant neoplasm of overlapping sites of brain
- C72.0 Malignant neoplasm of spinal cord
- C72.1 Malignant neoplasm of cauda equina
- C72.21 Malignant neoplasm of right olfactory nerve
- C72.22 Malignant neoplasm of left olfactory nerve
- C72.31 Malignant neoplasm of right optic nerve
- C72.32 Malignant neoplasm of left optic nerve
- C72.41 Malignant neoplasm of right acoustic nerve
- C72.42 Malignant neoplasm of left acoustic nerve
- C72.59 Malignant neoplasm of other cranial nerves
- C73 Malignant neoplasm of thyroid gland
- C75.0 Malignant neoplasm of parathyroid gland
- C75.1 Malignant neoplasm of pituitary gland
- C75.2 Malignant neoplasm of craniopharyngeal duct
- C75.3 Malignant neoplasm of pineal gland
- C75.4 Malignant neoplasm of carotid body
- C76.0 Malignant neoplasm of head, face and neck
- C77.0 Secondary and unspecified malignant neoplasm of lymph nodes of head, face and neck

- C79.31 Secondary malignant neoplasm of brain
- C83.31 Diffuse large B-cell lymphoma, lymph nodes of head, face, and neck
- C83.51 Lymphoblastic (diffuse) lymphoma, lymph nodes of head, face, and neck
- D32.0 Benign neoplasm of cerebral meninges
- D32.1 Benign neoplasm of spinal meninges
- D32.9 Benign neoplasm of meninges, unspecified
- D33.0 Benign neoplasm of brain, supratentorial
- D33.1 Benign neoplasm of brain, infratentorial
- D33.3 Benign neoplasm of cranial nerves
- D33.4 Benign neoplasm of spinal cord
- D33.7 Benign neoplasm of other specified parts of central nervous system
- D35.1 Benign neoplasm of parathyroid gland
- D35.2 Benign neoplasm of pituitary gland
- D35.3 Benign neoplasm of craniopharyngeal duct
- D35.4 Benign neoplasm of pineal gland
- D35.5 Benign neoplasm of carotid body
- D35.6 Benign neoplasm of aortic body and other paraganglia
- D40.0 Neoplasm of uncertain behavior of prostate
- D42.0 Neoplasm of uncertain behavior of cerebral meninges
- D42.1 Neoplasm of uncertain behavior of spinal meninges
- D43.0 Neoplasm of uncertain behavior of brain, supratentorial
- D43.1 Neoplasm of uncertain behavior of brain, infratentorial
- D43.4 Neoplasm of uncertain behavior of spinal cord
- D44.3 Neoplasm of uncertain behavior of pituitary gland
- D44.4 Neoplasm of uncertain behavior of craniopharyngeal duct
- D44.5 Neoplasm of uncertain behavior of pineal gland

REVISIONS

01-30-2009	<p>Policy first published on www.bcbsks.com.</p> <p>In policy section:</p> <ul style="list-style-type: none"> ▪ Added the following indications: <p>D. IMRT is considered medically necessary for treatment of <u>anal cancers</u>. E. IMRT may also be medically necessary for other radiosensitive tumors where critical structures cannot be adequately protected with standard 3D conformal radiotherapy. Medical necessity for the use of IMRT for these other indications will be considered individually and will require supporting records from the treating radiation oncologist including the dose volume histograms documenting the need for IMRT as opposed to conventional radiation therapy. The American Society for Therapeutic Radiology and Oncology (ASTRO) has a model policy which describes the indications for IMRT:</p>
	<p>"IMRT is not a replacement therapy for conventional or three-dimensional conformal radiation therapy methods. IMRT is considered reasonable and necessary in instances where sparing the surrounding normal tissue is of added benefit and at least one of the following conditions is met:</p> <ol style="list-style-type: none"> 1. The target volume is in close proximity to critical structures that must be protected. 2. The volume of interest must be covered with narrow margins to adequately protect immediately adjacent structures.

	<p>3. An immediately adjacent area has been previously irradiated and abutting portals must be established with high precision.</p> <p>4. The target volume is concave or convex, and critical normal tissues are within or around that convexity or concavity.</p> <p>5. Dose escalation is planned to deliver radiation doses in excess of those commonly utilized for similar tumor with conventional treatments."</p> <p>F. Other applications of IMRT are considered not medically necessary.</p>
	<ul style="list-style-type: none"> ▪ Added the following documentation information: <u>DOCUMENTATION</u> <p>As recommended by ASTRO, documentation in the patient's medical records must support:</p> <ol style="list-style-type: none"> 1. The reasonable and necessary requirements as outlined in the Policy section. 2. The prescription must define the dose to the target and the dose constraints to the nearby critical structures. 3. A note of medical necessity for IMRT, by the treating physician. 4. Signed IMRT inverse plan that meets prescribed dose constraints for the planning target volume (PTV) and surrounding normal tissue. 5. The target verification methodology must include the following: <ol style="list-style-type: none"> a. Documentation of the clinical treatment volume (CTV) and the planning target volume (PTV). b. Documentation of immobilization and patient positioning. 6. Independent basic dose calculations of monitor units have been performed for each beam before the patient's first treatment. 7. Documentation of fluence distributions (re-computed and measured in a phantom or dosimetry measuring device) is required. 8. Identification of structures that transverse high-and low-dose regions created by respiration is indicated. Voluntary breath-holding alone is not a satisfactory solution for accounting for organ motion.
	<p>In Coding section:</p> <ul style="list-style-type: none"> ▪ Reflected the applicable CPT codes 77300, 77301, 77332, 77333, 77334, 77418, 0073T
01-01-2010	<p>In Coding Section:</p> <ul style="list-style-type: none"> ▪ Added CPT Code: 77388
10-11-2011	<p>In the Policy section:</p> <ul style="list-style-type: none"> ▪ Item D, removed "anal cancers" and added "squamous cell carcinoma of the anus." ▪ Added Item " E. IMRT is considered medically necessary in the treatment of individuals with anaplastic thyroid cancer." ▪ Added Item "F. IMRT is considered medically necessary in individuals with pediatric tumors (e.g., Ewing Sarcoma, Wilms' Tumor)." ▪ Added Item "G. IMRT is considered medically necessary in individuals who require repeat irradiation of a field that has received prior irradiation." <p>Updated the Reference section.</p>
07-13-2012	<p>Updated the Description section.</p> <p>In the Policy section:</p> <ul style="list-style-type: none"> ▪ Added the following indication: "F. Carcinoma of the cervix." ▪ Added the following indication: "G. Whole pelvic radiotherapy for gynecologic malignancies." ▪ In Item J, inserted "3-D" to read "radiation oncologist including the 3-D dose

	<p>volume..."</p> <ul style="list-style-type: none"> ▪ In Item J, changed "histograms" to "histogram" ▪ In Item J, removed "as opposed to conventional radiation therapy" from the end of paragraph. <p>In the Documentation section:</p> <ul style="list-style-type: none"> ▪ Revised the following language: "As recommended by ASTRO, documentation in the patient's medical records must support: <ol style="list-style-type: none"> 1. The reasonable and necessary requirements as outlined in the Policy section. 2. The prescription must define the dose to the target and the dose constraints to the nearby critical structures. 3. A note of medical necessity for IMRT, by the treating physician. 4. Signed IMRT inverse plan that meets prescribed dose constraints for the planning target volume (PTV) and surrounding normal tissue. 5. The target verification methodology must include the following: <ol style="list-style-type: none"> a. Documentation of the clinical treatment volume (CTV) and the planning target volume (PTV). b. Documentation of immobilization and patient positioning. 6. Independent basic dose calculations of monitor units have been performed for each beam before the patient's first treatment. 7. Documentation of fluence distributions (re-computed and measured in a phantom or dosimetry measuring device) is required. 8. Identification of structures that transverse high-and low-dose regions created by respiration is indicated. Voluntary breath-holding alone is not a satisfactory solution for accounting for organ motion."
	<p>In the Coding section:</p> <ul style="list-style-type: none"> ▪ Added the following Diagnosis codes: 180.0-180.9, 182.0-182.8, V10.40, V10.41
	<p>Updated Reference section.</p>
<p>10-15-2012</p>	<p>In the Policy section:</p> <ul style="list-style-type: none"> ▪ Added statement, "IMRT is considered medically necessary for the following indications:" to the beginning of the policy section. ▪ In Item A, removed "IMRT of the prostate is considered medically necessary in patients with non-metastatic..." and added "of the prostate" to read "Prostate cancer for dose escalation >75Gy of the prostate and for..." ▪ In Item B, removed, "IMRT is considered medically necessary in the treatment of patients with" to read "head and neck cancer, with the exception..." ▪ In item C, removed "IMRT is considered medically necessary in patients with CNS lesions" and added "Cancer involving the CNS" ▪ In Item D, removed IMRT is considered medically necessary for patients of squamous cell" and added "or vulva" to read "Carcinoma of the anus and vulva." ▪ In Item E, removed "IMRT is considered medically necessary in the treatment of individuals with" to read "Anaplastic thyroid cancer." ▪ In Item H, removed "IMRT is considered medically necessary in individuals with" to read "Pediatric tumors (e.g., Ewing Sarcoma, Wilms' Tumor)." ▪ Added Item I, "Breast cancer when at least one of the following is met: <ol style="list-style-type: none"> 1. Heart—3D results in $\geq 25\%$ of heart receiving $\geq 30\text{Gy}$; <p style="text-align: center;">OR</p>

	<p>2. Lung—3D results in $\geq 30\%$ of ipsilateral lung receiving ≥ 20 Gy, OR 3D results in $\geq 20\%$ of combined lung volume receiving ≥ 20 Gy; OR</p> <p>3. Skin / Soft Tissue—3D results in $\geq 5\%$ of intended breast receiving $\geq 7\%$ of prescribed dose OR Medial lesion where 3D results in $\geq 10\%$ of contralateral breast receiving ≥ 10 Gy."</p> <p>▪ Added Item J, "Esophagus, Stomach, Pancreas, Hepatobiliary Tract, Rectum, Colon, Small Bowel, when at least one of the following is met:</p> <p>1. Heart—3D result in $\geq 50\%$ of heart receiving ≥ 30 Gy, OR</p> <p>2. Lung—3D results in $\geq 30\%$ of combined lung volume receiving $\geq 20\%$ Gy, OR Mean lung dose ≥ 20 Gy; OR</p> <p>3. Spinal Cord—3D result in any portion of the spinal cord receiving a dose above 45 Gy; OR</p> <p>4. Liver—3D results in $\geq 60\%$ of liver volume receiving ≥ 30 Gy, OR Mean liver dose ≥ 32 Gy; OR</p> <p>5. Kidney—3D results in $\geq 33\%$ of combined kidney volume receiving ≥ 20 Gy (two functional kidneys are present); OR For one functioning kidney or kidney transplant, IMRT provides a lower dose than achievable with 3D; OR</p> <p>6. Small Intestine—3D results in ≥ 195cc of small intestine receiving ≥ 45 Gy; OR</p> <p>7. Stomach—3D results in $\geq 10\%$ of stomach receiving ≥ 45 Gy OR 5% receiving ≥ 50 Gy; OR</p> <p>8. Femoral head—3D results in a femoral head receiving ≥ 45 Gy."</p> <p>▪ Added Item K, "Lung, when at least one of the following is met:</p> <p>1. Heart—3D results in $\geq 50\%$ of heart receiving ≥ 30 Gy; OR</p> <p>2. Lung—3D result in $\geq 30\%$ of non-cancerous combined lung volume receiving ≥ 20 Gy"</p> <p>▪ Added Item L, "Lymphomas or Sarcomas of Retroperitoneum, Chest Wall and Thorax, when at least one of the following is met:</p> <p>1. Heart—3D results in $\geq 50\%$ of heart receiving ≥ 30 Gy; OR</p> <p>2. Lung—3D results in $\geq 30\%$ of combined lung volume receiving \geq</p>
--	--

	<p>20Gy OR Mean lung dose of ≥ 20 Gy; OR</p> <p>3. Spinal cord—3D results in any portion of the spinal cord receiving a dose above 45 Gy OR</p> <p>4. Liver—3D results in 60% of liver volume receiving ≥ 30 Gy OR Mean liver dose ≥ 32 Gy; OR</p> <p>5. Femoral head—3D results in a femoral head receiving ≥ 45 Gy; OR</p> <p>6. Small intestine—3D results in ≥ 195cc of small intestine receiving ≥ 45 Gy; OR</p> <p>7. Stomach—3D results in $\geq 10\%$ of stomach receiving ≥ 45 Gy OR 5% receiving ≥ 50 Gy; OR</p> <p>8. Rectosigmoid—3D results in $\geq 60\%$ of rectosigmoid area receiving ≥ 30 Gy; OR</p> <p>9. Bladder—3D results in $\geq 35\%$ of bladder receiving ≥ 45 Gy; OR</p> <p>10. Kidney—3D results in 33% of combined kidney volume receiving ≥ 20 Gy (two functional kidneys are present) OR For one functioning kidney or kidney transplant IMRT provides a lower dose than achievable with 3D."</p> <ul style="list-style-type: none"> ▪ Added Item M, "Sarcomas of the Extremities, when at least one of the following is met: <ul style="list-style-type: none"> 1. Head / Neck—IMRT covered if head and neck structures would receive any radiation via 3D; OR 2. Femur—3D results in $\geq 50\%$ of contiguous femur cortex receiving ≥ 50 Gy" ▪ In Item N, removed "IMRT is considered medically necessary in" to read "Individuals who require repeat irradiation..." ▪ In Item O, removed "IMRT may also be medically necessary for" to read "Other radiosensitive tumors where critical structures..."
	<p>In the Coding section:</p> <ul style="list-style-type: none"> ▪ Added Diagnosis codes: 174.0-174.9
12-11-2013	Policy reviewed.
	<p>In Coding section:</p> <ul style="list-style-type: none"> ▪ Added ICD-10 Diagnosis codes. <i>(Effective October 1, 2014)</i>
	Updated Reference section.

REFERENCES

1. Donovan E, Bleakley N, Denholm E et al. Randomised trial of standard 2D radiotherapy (RT) versus intensity modulated radiotherapy (IMRT) in patients prescribed breast radiotherapy. *Radiother Oncol* 2007; 82(3):254-64.
2. Pignol JP, Olivetto I, Rakovitch E et al. A multicenter randomized trial of breast intensity-modulated radiation therapy to reduce acute radiation dermatitis. *J Clin Oncol* 2008; 26(13):2085-92.
3. Violet JA, Harmer C. Breast cancer: improving outcome following adjuvant radiotherapy. *Br J Radiol* 2004; 77(922):811-20.
4. Arthur DW, Morris MM, Vicini FA. Breast cancer: new radiation treatment options. *Oncology (Williston Park)* 2004; 18(13):1621-9; discussion 29-30, 36-38.
5. Coles CE, Moody AM, Wilson CB et al. Reduction of radiotherapy-induced late complications in early breast cancer: the role of intensity-modulated radiation therapy and partial breast irradiation. Part II--Radiotherapy strategies to reduce radiation-induced late effects. *Clin Oncol (R Coll Radiol)* 2005; 17(2):98-110.
6. Formenti SC, Truong MT, Goldberg JD et al. Prone accelerated partial breast irradiation after breast-conserving surgery: preliminary clinical results and dose-volume histogram analysis. *Int J Radiat Oncol Biol Phys* 2004; 60(2):493-504.
7. Alonso-Basanta M MS, Lymberis S et al. Dosimetric comparisons of supine versus prone radiation: implications on normal tissue toxicity. *Int J Radiat Oncol Biol Phys* 2005; 63(2 suppl1):S182-3.
8. Remouchamps VM, Vicini FA, Sharpe MB et al. Significant reductions in heart and lung doses using deep inspiration breath hold with active breathing control and intensity-modulated radiation therapy for patients treated with locoregional breast irradiation. *Int J Radiat Oncol Biol Phys* 2003; 55(2):392-406.
9. Frazier RC, Vicini FA, Sharpe MB et al. Impact of breathing motion on whole breast radiotherapy: a dosimetric analysis using active breathing control. *Int J Radiat Oncol Biol Phys* 2004; 58(4):1041-7.
10. Chang JY, Liu HH, Komaki R. Intensity modulated radiation therapy and proton radiotherapy for non-small cell lung cancer. *Curr Oncol Rep* 2005; 7(4):255-9.
11. Dayes I, Rumble RB, Bowen J et al. Intensity-modulated radiotherapy in the treatment of breast cancer. *Clin Oncol (R Coll Radiol)* 2012; 24(7):488-98.
12. Staffurth J. A review of the clinical evidence for intensity-modulated radiotherapy. *Clin Oncol (R Coll Radiol)* 2010; 22(8):643-57.
13. Donovan EM, Bleackley NJ, Evans PM et al. Dose-position and dose-volume histogram analysis of standard wedged and intensity modulated treatments in breast radiotherapy. *Br J Radiol* 2002; 75(900):967-73.
14. Donovan EM, Yarnold JR, Adams EJ et al. An investigation into methods of IMRT planning applied to breast radiotherapy. *Br J Radiol* 2008; 81(964):311-22.
15. Barnett GC, Wilkinson J, Moody AM et al. A randomised controlled trial of forward-planned radiotherapy (IMRT) for early breast cancer: baseline characteristics and dosimetry results. *Radiother Oncol* 2009; 92(1):34-41.
16. Barnett GC, Wilkinson JS, Moody AM et al. Randomized controlled trial of forward-planned intensity modulated radiotherapy for early breast cancer: interim results at 2 years. *Int J Radiat Oncol Biol Phys* 2012; 82(2):715-23.
17. McDonald MW, Godette KD, Butker EK et al. Long-term outcomes of IMRT for breast cancer: a single-institution cohort analysis. *Int J Radiat Oncol Biol Phys* 2008; 72(4):1031-40.

18. Kestin LL SM, Frazier RC et al. Intensity modulation to improve dose uniformity with tangential breast radiotherapy: Initial clinical experience. *Int J Radiat Oncol Biol Phys* 2005; 48(5):1559-68.
19. Vicini FA, Sharpe M, Kestin L et al. Optimizing breast cancer treatment efficacy with intensity-modulated radiotherapy. *Int J Radiat Oncol Biol Phys* 2002; 54(5):1336-44.
20. Selvaraj RN, Beriwal S, Pourarian RJ et al. Clinical implementation of tangential field intensity modulated radiation therapy (IMRT) using sliding window technique and dosimetric comparison with 3D conformal therapy (3DCRT) in breast cancer. *Med Dosim* 2007; 32(4):299-304.
21. Hardee ME, Raza S, Becker SJ et al. Prone hypofractionated whole-breast radiotherapy without a boost to the tumor bed: comparable toxicity of IMRT versus a 3D conformal technique. *Int J Radiat Oncol Biol Phys* 2012; 82(3):e415-23.
22. Freedman GM, Li T, Nicolaou N et al. Breast intensity-modulated radiation therapy reduces time spent with acute dermatitis for women of all breast sizes during radiation. *Int J Radiat Oncol Biol Phys* 2009; 74(3):689-94.
23. Keller LM, Sopka DM, Li T et al. Five-year results of whole breast intensity modulated radiation therapy for the treatment of early stage breast cancer: the Fox Chase Cancer Center experience. *Int J Radiat Oncol Biol Phys* 2012; 84(4):881-7.
24. Coon AB, Dickler A, Kirk MC et al. Tomotherapy and multifield intensity-modulated radiotherapy planning reduce cardiac doses in left-sided breast cancer patients with unfavorable cardiac anatomy. *Int J Radiat Oncol Biol Phys* 2010; 78(1):104-10.
25. Leonard C, Carter D, Kercher J et al. Prospective trial of accelerated partial breast intensity-modulated radiotherapy. *Int J Radiat Oncol Biol Phys* 2007; 67(5):1291-8.
26. Livi L, Buonamici FB, Simontacchi G et al. Accelerated partial breast irradiation with IMRT: new technical approach and interim analysis of acute toxicity in a phase III randomized clinical trial. *Int J Radiat Oncol Biol Phys* 2010; 77(2):509-15.
27. Rudat V, Alaradi AA, Mohamed A et al. Tangential beam IMRT versus tangential beam 3D-CRT of the chest wall in postmastectomy breast cancer patients: a dosimetric comparison. *Radiat Oncol* 2011; 6:26.
28. Bezjak A, Rumble RB, Rodrigues G et al. Intensity-modulated radiotherapy in the treatment of lung cancer. *Clin Oncol (R Coll Radiol)* 2012; 24(7):508-20.
29. Liao ZX, Komaki RR, Thames HD, Jr. et al. Influence of technologic advances on outcomes in patients with unresectable, locally advanced non-small-cell lung cancer receiving concomitant chemoradiotherapy. *Int J Radiat Oncol Biol Phys* 2010; 76(3):775-81.
30. Holloway CL, Robinson D, Murray B et al. Results of a phase I study to dose escalate using intensity modulated radiotherapy guided by combined PET/CT imaging with induction chemotherapy for patients with non-small cell lung cancer. *Radiother Oncol* 2004; 73(3):285-7.
31. Sura S, Gupta V, Yorke E et al. Intensity-modulated radiation therapy (IMRT) for inoperable non-small cell lung cancer: the Memorial Sloan-Kettering Cancer Center (MSKCC) experience. *Radiother Oncol* 2008; 87(1):17-23.
32. Jiang ZQ, Yang K, Komaki R et al. Long-term clinical outcome of intensity-modulated radiotherapy for inoperable non-small cell lung cancer: the MD Anderson experience. *Int J Radiat Oncol Biol Phys* 2012; 83(1):332-9.
33. Kwint M, Uyterlinde W, Nijkamp J et al. Acute esophagus toxicity in lung cancer patients after intensity modulated radiation therapy and concurrent chemotherapy. *Int J Radiat Oncol Biol Phys* 2012; 84(2):e223-8.
34. Govaert SL, Troost EG, Schuurbiens OC et al. Treatment outcome and toxicity of intensity-modulated (chemo) radiotherapy in stage III non-small cell lung cancer patients. *Radiat Oncol* 2012; 7:150.

35. National Comprehensive Cancer Network (NCCN). Breast Cancer. National Comprehensive Cancer Network Clinical Practice Guidelines in Oncology. V.2.2013. Available online at: http://www.nccn.org/professionals/physician_gls/pdf/breast.pdf. Last accessed March 22, 2013.
36. National Comprehensive Cancer Network (NCCN). Non-small Cell Lung Cancer. National Comprehensive Cancer Network Clinical Practice Guidelines in Oncology. V.2.2013. Available online at: http://www.nccn.org/professionals/physician_gls/pdf/nscl.pdf. Last accessed March 22, 2013.
37. National Comprehensive Cancer Network (NCCN). Small Cell Lung Cancer. National Comprehensive Cancer Network Clinical Practice Guidelines in Oncology. V.2.2013. Available online at: http://www.nccn.org/professionals/physician_gls/pdf/sclc.pdf. Last accessed March 22, 2013.
38. Marks LB, Bentzen SM, Deasy JO et al. Radiation dose-volume effects in the lung. *Int J Radiat Oncol Biol Phys* 2010; 76(3 Suppl):S70-6.
39. American College of Radiology (ACR). ACR Practice Guideline for Intensity Modulated Radiation Therapy. 2002 (Res. 17). ACR Practice Guideline. Reston, VA: ACR; effective January 1, 2003:705-710. Available at: http://www.acr.org/s_acr/bin.asp?TrackID=&SID=1&DID=12234&CID=1075&VID=2&DOC=File.PDF
40. American Society for Therapeutic Radiology and Oncology (ASTRO). The ASTRO/ACR Guide to Radiation Oncology Coding. 2007; Fairfax, VA. ASTRO; 2007. Available at: <http://www.astro.org/publications>.
41. Blue Cross Blue Shield Association. Special Report: Intensity Modulation Radiation Therapy for Cancer of the Breast or Lung. *TEC Assessment*, 2005; 20(13).
42. National Comprehensive Cancer Network (NCCN). Clinical guidelines in Oncology. Breast cancer v.2.2007. Rockledge, PA: NCCN; 2007. Available at http://www.nccn.org/professionals/physician_gls/PDF/breast/pdf.
43. American Society for Therapeutic Radiology and Oncology (ASTRO) Model Policy on Intensity Modulated Radiation Therapy (IMRT) 03-2007.

Other References

1. Blue Cross and Blue Shield of Kansas Radiology Liaison Committee, February 2008; February 2009; February 2010; February 2011.
2. Blue Cross and Blue Shield of Kansas Radiology Liaison Committee, Consent Ballot, January 2009.
3. Blue Cross and Blue Shield of Kansas Medical Consultant, Practicing Board-Certified Radiation Oncologist (548), September 2011.
4. Blue Cross and Blue Shield of Kansas Medical Consultant, Practicing Board Certified Radiation Oncologist (548), March 2012.
5. Blue Cross and Blue Shield of Kansas Medical Consultant, Practicing Board Certified Radiation Oncologist (548), April 2012.