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## Subject: Image-Guided Radiation Therapy

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<a href="#">Position Statement</a>	<a href="#">Billing/Coding</a>	<a href="#">Reimbursement</a>	<a href="#">Program Exceptions</a>	<a href="#">Definitions</a>	<a href="#">Related Guidelines</a>
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### DESCRIPTION:

Image-guided radiation therapy (IGRT) is radiation therapy that employs imaging to maximize accuracy and precision throughout the entire process of treatment delivery. This process can include target and normal tissue delineation, radiation delivery, and adaptation of therapy to anatomic and biological and positional changes over time in individuals. IGRT is the use of imaging at the time of treatment delivery to ensure that the location of the target relative to the treatment beams based on a predetermined plan is reliably reproduced. IGRT is applicable to highly conformal treatment modalities, such as 3-D conformal radiation therapy (CRT), intensity-modulated radiation therapy (IMRT), or heavy particle therapy (e.g., proton). IGRT is considered an integral component of treatment delivery with stereotactic body radiation therapy (SBRT) or stereotactic ablative radiotherapy (SABR). At the time of treatment delivery, IGRT is employed to determine the location of the target (and often the surrounding normal organs) at some predetermined frequency. The target location may be determined by a range of methods, from soft-tissue volumetric imaging (e.g., CT, ultrasound, MRI) to localization of surrogates, such as bone, implanted fiducial markers or external surface markers or features (e.g., by planar imaging or fluoroscopy, electromagnetic localization or optical surface imaging). The match or discrepancy between the simulated location and the “live” IGRT measurement prior to treatment delivery may be determined manually (e.g., visual alignment of the two image datasets), or in some cases, by using automated image analysis software. If a discrepancy is found, a correction is applied. In this manner, the treatment will be delivered precisely and accurately according to the treatment plan approved by the radiation oncologist. (ACR-ASTRO, 2019)

Several systems, devices, and components for IGRT have received U.S. Food and Drug Administration (FDA) 510(k) clearance (e.g., Calypso® 4D localization System, RPM Respiratory Gating System).

## POSITION STATEMENT:

Image-guided radiation therapy (IGRT) (any modality) **meets the definition of medical necessity** for the following indications:

- Bony anatomy fails to delineate tumor location
- Dose escalation is planned beyond the usual doses for similar tumors
- During radiation therapy (e.g., intensity-modulated radiation therapy (IMRT), proton beam therapy, stereotactic body radiation therapy (SBRT))
- Implanted fiducial markers
- Reduction of radiation dose to sensitive normal structures (e.g., left-sided breast radiation therapy with deep inspiration breath hold technique (DIBH) to spare heart radiation exposure)
- Target volume is in close proximity to critical structures that must be protected
- Target volume located near or within critical structures and/or in tissue with inherent setup variation
- Target volume that is subject to daily variation that is due to internal motion
- Target where the adjacent area has been previously irradiated and abutting fields must be precise
- Volume of interest that must be covered with narrow margins to adequately protect immediately adjacent structures.

## BILLING/CODING INFORMATION:

### CPT Coding:

77014	Computed tomography guidance for placement of radiation therapy fields
77387	Guidance for localization of target volume for delivery of radiation treatment, includes intrafraction tracking, when performed

### HCPCS Coding:

G6001	Ultrasonic guidance for placement of radiation therapy fields
G6002	Stereoscopic X-ray guidance for localization of target volume for the delivery of radiation therapy
G6017	Intra-fraction localization and tracking of target or patient motion during delivery of radiation therapy (e.g., 3D positional tracking, gating, 3D surface tracking), each fraction of treatment

### LOINC Codes:

The following information may be required documentation to support medical necessity: physician history and physical, physician progress notes, plan of treatment and reason for imaged-guided radiation therapy (IGRT).

Documentation Table	LOINC Codes	LOINC Time Frame	LOINC Time Frame Modifier Codes Narrative
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		<b>Modifier Code</b>	
Physician history and physical	28626-0	18805-2	Include all data of the selected type that represents observations made six months or fewer before starting date of service for the claim
Attending physician progress note	18741-9	18805-2	Include all data of the selected type that represents observations made six months or fewer before starting date of service for the claim
Plan of treatment	18776-5	18805-2	Include all data of the selected type that represents observations made six months or fewer before starting date of service for the claim

## REIMBURSEMENT INFORMATION:

Refer to section entitled [POSITION STATEMENT](#).

## PROGRAM EXCEPTIONS:

**Federal Employee Program (FEP):** Follow FEP guidelines.

**State Account Organization (SAO):** Follow SAO guidelines.

### Medicare Advantage products:

No National Coverage Determination (NCD) and/or Local Coverage Determination (LCD) were found at the time of the last guideline review date.

## DEFINITIONS:

None applicable.

## RELATED GUIDELINES:

None applicable.

## OTHER:

Other names used to report image-guided radiation therapy (IGRT):

Dynamic Targeting IGRT  
Electromagnetic Transponders  
Intra-fraction Image Guidance  
On-Board Imager  
Real-Time Intra-Fraction

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## COMMITTEE APPROVAL:

This Medical Coverage Guideline (MCG) was approved by the Florida Blue Medical Policy and Coverage Committee on 04/27/23.

## GUIDELINE UPDATE INFORMATION:

02/15/10	New Medical Coverage Guideline.
01/01/11	Annual HCPCS update; revised 55876 code descriptor.

02/15/11	Annual review. Maintain position statement. Updated references.
10/01/11	Revision; formatting changes.
05/15/14	Annual review. Maintain position statement. Added heading for “intra-fraction, description section. Revised FDA statement for Calypso 4D Localization System, description section. Added Medicare Advantage products program exception. Updated references.
01/01/15	Annual HCPCS code update. Deleted 0197T, 76050 and 77421. Added 77387, G6001, G6002 and G6017.
05/15/15	Annual review; position statement unchanged. Updated description and references.
05/01/16	Revision; deleted “for Treatment Planning and Delivery” from MCG name; deleted position statement for intra-fraction localization and tracking of target and imaging modalities; added position statement for image-guided radiation therapy; deleted 32553, 49411 and 55876; added LOINC codes; updated program exception; updated references.
11/15/16	Revision; revised position statement. Updated references.
10/15/17	Revision; revised position statement. Updated references.
02/15/18	Revision; updated position statement and definitions.
01/01/19	Annual HCPCS code update. Revised 77387 code descriptor.
03/15/21	Review/update. Maintain position statement. Updated references.
05/15/23	Review: revised description and position statement. Updated references.