

National Imaging Associates, Inc.*	
Clinical guidelines	Original Date: January 2015
LOW DOSE CT FOR LUNG CANCER SCREENING	
CPT Codes: 71271	Last Revised Date: November April 20219
Guideline Number: NIA_CG_020-1	Implementation Date: January 20221

INDICATIONS FOR LOW_DOSE CT (LDCT) FOR LUNG CANCER SCREENING (LDCT):

For Annual Lung Cancer Screening:

The use of low-dose, non-contrast spiral (helical) multi-detector CT imaging as a screening technique for lung cancer is considered **medically necessary ONLY** when used to screen for lung cancer for certain high-risk, **asymptomatic** individuals, i.e., no acute <u>lung-lung-</u>related symptoms, when **ALL** of the following criteria are met (<u>Mazzone, 2018; NCCN, **2019**USPSTF, 2021):</u>

Group 1: Group 1:

- Individual is between 5<u>0</u>5-80 years of age*; AND
- There is at least a 230 pack-year history of cigarette smoking; AND
- If the individual is a former smoker, that individual had quit smoking within the previous 15 years.
- *May approve for individuals over the age limit if the individual is a candidate for and willing to undergo curative treatment

Group 2:

<u>Yearly Low-Dose CT surveillance after completion of definitive treatment of non-small cell lung</u> cancer as per these parameters (NCCN version 4.2021):

- Stage I-II (treated with surgery +/- chemotherapy)
 - starts at year 2-3 of surveillance)
- Stage I-II (treated primarily with radiation) or stage III-IV with all sites treated with definitive intent
 - starts at year 5 of surveillance

^{*} National Imaging Associates, Inc. (NIA) is a subsidiary of Magellan Healthcare, Inc.

^{1—}CT Lung CA Screening

Group 2:

- Age ≥ 50 years old; AND
- ≥ 20 pack-year history of smoking; AND
- Additional risk factors (other than second hand smoke)*

*Additional risk factors include: Survivors of lung cancer, lymphoma, cancers of the head and neck and bladder (smoking related cancers), first degree family members with a history of lung cancer, history of COPD or pulmonary fibrosis, radon exposure, retinoblastoma, Li Fraumeni syndrome, occupational exposure to arsenic, chromium, asbestos, nickel, cadmium, beryllium, silica, diesel fumes, coal smoke and soot.

Nodule on initial LDCT (Follow-up low dose CT is approvable): (Wood, 2018)

- <u>Table 1</u> shows the follow-up interval at which LDCT can be approved to reduce radiation dose (ACR, 2019)
- If multiple nodules, the largest and type is used for decision

Table 1: Lung-RADS® Assessment Categories (ACR, 2019)

Category	Lung- RADS	Findings	Management
Descriptor	Score	rindings	management
Incomplete	0	Prior chest CT examination(s) being located for comparison	Additional lung cancer screening CT images and/or
		Part or all of lungs cannot be evaluated	comparison to prior chest CT examinations is needed
Negative		No lung nodules	
No nodules and	1	Nodule(s) with specific calcifications: complete, central, popcorn, concentric	
definitely benign		rings and fat containing nodules	
nodules		Perifissural nodule(s) (See Footnote 11)	
		< 10 mm (524 mm³)	
Benign Appearance or		Solid nodule(s): < 6 mm (< 113 mm ³)	Continue annual
Behavior		new < 4 mm (< 34 mm ³)	screening with LDCT in
Nodules with a very low		Part solid nodule(s): < 6 mm total diameter (< 113 mm ³) on	12 months
likelihood of becoming a	2	baseline screening	
clinically active cancer		Non solid nodule(s) (GGN):	
due to size or lack of growth		<30 mm (<14137 mm³) OR ≥ 30 mm (≥ 14137 mm³) and unchanged	
g. 2.1.2.		or slowly growing	
		Category 3 or 4 nodules unchanged for ≥ 3 months	
		Solid nodule(s):	
Probably Benign		≥ 6 to < 8 mm (≥ 113 to < 268 mm³) at baseline OR	
Probably benign		new 4 mm to < 6 mm (34 to < 113 mm ³)	
finding(s) - short term	3	Part solid nodule(s)	6 month LDCT
follow up suggested; includes nodules with a	3	≥ 6 mm total diameter (≥ 113 mm³) with solid component < 6 mm (< 113 mm³) OR	6 month LDC1
low likelihood of		new < 6 mm total diameter (< 113 mm ³)	
becoming a clinically active cancer		Non solid nodule(s) (GGN) ≥ 30 mm (≥ 14137 mm ³) on	
		baseline CT or new	
		Solid nodule(s):	
		≥ 8 to < 15 mm (≥ 268 to < 1767 mm³) at baseline OR	
Suspicious		growing < 8 mm (< 268 mm³) OR	
Suspicious		new 6 to < 8 mm (113 to < 268 mm ³) Part solid nodule(s):	3 month LDCT; PET/CT may be
Findings for which	4A	≥ 6 mm (≥ 113 mm³) with solid	used when there is a ≥ 8 mm (≥ 268 mm³) solid component
additional diagnostic testing is recommended		component ≥ 6 mm to < 8 mm (≥ 113 to < 268 mm³) OR	266 mm²) solid component
•		with a new or growing < 4 mm (< 34 mm ³)	
		solid component	
		Endobronchial nodule Solid nodule(s)	Chest CT with or without
		≥ 15 mm (≥ 1767 mm ³) OR	contrast, PET/CT and/or tissue
Very Suspicious		new or growing, and ≥ 8 mm (≥ 268 mm³)	sampling depending on the *probability of malignancy and
very ouspicious	4B	Part solid nodule(s) with: a solid component ≥ 8 mm (≥ 268 mm ³)	comorbidities. PET/CT may be
Findings for which		OR	used when there is a ≥ 8 mm (≥ 268 mm³) solid component.
additional diagnostic testing and/or tissue		a new or growing ≥ 4 mm (≥ 34 mm³)	(≥ 268 mm²) solid component. For new large nodules that
sampling is recommended		solid component	develop on an annual repeat
recommended	4X	Category 3 or 4 nodules with additional features or imaging findings that	screening CT, a 1 month LDCT may be recommended to
	4,7	increases the suspicion of malignancy	address potentially infectious or inflammatory conditions
Other			or initialitinatory conditions
Clinically Significant or Potentially Clinically	s	Modifier - may add on to category 0-4	As appropriate to the specific
Significant Findings	3	coding	finding
(non lung cancer)			

BACKGROUND:

Smoking-related lung cancer is the leading cause of cancer deaths in both men and women in the United States. Treatment for most lung cancer is focused on surgery which is usually curative only when the tumors are very small. Screening for early lung cancer with sputum cytology and chest x-rays has not been successful in reducing deaths from lung cancer. However, in 2011, a large, prospective, multicenter trial was published that showed CT Chest screening identified early cancers better than other approaches and reduced the death rate from lung cancer. In 2014, the United States Preventive Service Task Force (USPSTF) recommended annual low-low-dose CT Chest screening (CPT® code 71271) for people with current or recent past smoking histories.

All screening and follow-up chest CT scans to be performed at low dose (100-120 kVp and 40-60 mAs), unless evaluating mediastinal findings or lymph nodes, where standard dose CT with IV contrast may be more appropriate (NCCN, 20192021).

OVERVIEW:

Screening should be discontinued once a person has not smoked for 15 years or develops a health problem that substantially limits life expectancy or the ability or willingness to have curative lung surgery.

POLICY HISTORY:

<u>Date</u>	Summary	
April 2021	 Added data about expanding screening to older patients that are willing to have and that are candidates for definitive treatment for lung cancer, based on NCCN recommendations Added long term surveillance in patients who received definitive treatment for non small cell lung cancer 	
March 10, 2021	 Eliminated groupings (group 1 and group 2) for lung cancer screening and changed age of 55-80 years to 50-80 years; removed 30changed to 20 pack year history of cigarette smoking and requirement of additional risk factors (USPSTF 2021) 	
November 9, 2020	Replaced CPT code G0297 with 71271	
May 2020	 Lung Cancer Screening: Changed upper age limit from 77 to 80 yrs old Added: Age ≥ 50 years old; AND 	

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	 ≥ 20 pack-year history of smoking; AND 		
	 Additional risk factors (other than second-hand 		
	smoke)*		
	*Additional risk factors include: Survivors of lung cancer, lymphoma,		
	cancers of the head and neck and bladder (smoking related cancers),		
	first degree family members with a history of lung cancer, history of		
	COPD or pulmonary fibrosis, radon exposure, retinoblastoma, Li		
	Fraumeni syndrome, occupational exposure to arsenic, chromium,		
	asbestos, nickel, cadmium, beryllium, silica, diesel fumes, coal smoke		
	and soot		
	 Updated the follow-up interval for LDCT information, using the 		
	ACR 2019 Lung RADS chart		
	Updated background information		
May 2019	Criteria for repeating at less than one year were added.		
	Upper age range changed from 80 to 77 years of age		
	Chart added for the f/u interval at which LDCT can be approved		
	to reduce radiation dose		

Review Date: May 2019 Review Summary:

- Criteria for repeating at less than one year were added.
- Upper age range changed from 80 to 77 years of age
- Chart added for the f/u interval at which LDCT can be approved to reduce radiation dose

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- Lung Cancer Screening:
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- Updated the follow-up interval for LDCT information, using the ACR 2019 Lung RADS chart
- Updated background information

Review Date: November 9, 2020

Review Summary: Replaced CPT code G0297 with 71271

REFERENCES:

American College of Radiology (ACR). Lung - RADS Assessment Categories v1.1. 2019. https://www.acr.org/Clinical-Resources/Reporting-and-Data-Systems/Lung-Rads.

Mazzone PJ, Silvestri GA, Patel S, et al. Screening for lung cancer CHEST guideline and expert panel report. *Chest.* 2018; 153(4):954-985.

National Comprehensive Cancer Network (NCCN). NCCN Guidelines Version <u>4</u>1.202<u>1</u>9 – <u>Lung Cancer ScreeningNon-Small Cell Lung Cancer</u>. <u>May 14, 2019March 3, 2021</u>. https://www.nccn.org/professionals/physician gls/pdf/nscl.pdf.

U.S. Preventive Services Task Force (USPSTF). Lung Cancer: Screening. Final recommendation statement, March 9, 2021.

https://uspreventiveservicestaskforce.org/uspstf/recommendation/lung-cancer-screening.

Wood DE, Kazerooni EA, Baum SL, et al. Clinical practice guidelines in oncology: Lung cancer screening. Version 3.2018. *J Natl Compr Canc Netw.* 2018; 16(4):412–441.

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GENERAL INFORMATION

It is an expectation that all patients receive care/services from a licensed clinician. All appropriate supporting documentation, including recent pertinent office visit notes, laboratory data, and results of any special testing must be provided. If applicable: All prior relevant imaging results and the reason that alternative imaging cannot be performed must be included in the documentation submitted.

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