

## Talking Points to Address the Annals Article: Rates of Downstream Procedures and Complications Associated with Lung Cancer Screening in Routine Clinical Practice — A Retrospective Cohort Study

Lung cancer screening (LCS) is a vital tool for early detection, enhancing survival rates.<sup>1</sup> Performed using a low-dose CT (LDCT) scan, LCS identifies potential issues before symptoms arise, enabling timely intervention and improved outcomes.<sup>2</sup> The American College of Radiology® (ACR®) has adopted a unified coding system for grading lung nodules; Lung-RADS®, now in its third revision, provides specific guidelines for next steps when a nodule is deemed suspicious. Diagnostic studies may include biopsy through bronchoscopy, EBUS, CT guided IR and surgery.<sup>3</sup> The study described in the publication addresses complications related to biopsy and resection procedures. The information presented can be misleading, instill fear among physicians, and discourage other physicians from participating in the lung cancer screening and associated follow-up process when abnormal findings are present. Below are talking points suggested when asked about the article published.

- The PROSPR (Populations-Based Research to Optimize the Screening Process) researchers studied 9,266 individuals who received a baseline LDCT lung screening completed between 2014 to 2018; the data presented is at least five years old. Lung cancer screening has evolved since 2018 and quality indicators show fewer harms than in this study.
- The false positive rate (FPR) for LCS is quoted in the article as being high at 73.4% for National Lung Screening Trial (NLST) after initial screening.<sup>4</sup> The 73.4% reported was the false discovery rate (FDR) for NLST, NOT the FPR. Using the FDR as an indicator for complications is disingenuous as the NLST predated Lung-RADS.<sup>3</sup> In NLST the 96.4% overall FPR (17,497/18,146) is really the FDR.<sup>4</sup> The overall FPR for the NLST (pre-Lung-RADS) was 23.3% (18,146 - 649/75,126). The current FPR for lung cancer screening are less than 10.4%<sup>5</sup>

LCS False Positive is not ~~96.4%~~ as frequently reported

	Cancer	No Cancer	
Screen +ve	True + (A)	False + (B)	A + B
Screen -ve	False - (C)	True - (D)	C + D
	A + C	B + D	

False Positive Rate is  $B / (B + D)$

**FALSE POSITIVE** 1. A positive test result in a person who does not possess the attribute (e.g., the disease) for which the test is conducted. The labeling of a healthy person as diseased in SCREENING. A Dictionary of Epidemiology, Oxford University Press.

- PROSPR had higher rates of current smoking (55% vs. 48%) and COPD (35% vs. 18%) than found in the NLST population.



- The PROSPR population was significantly older than the NLST population, with 52% of individuals 65 and older, versus 26.6% in NLST. In the article, they compare patients with abnormal findings on screening and subsequent invasive procedures' (59.4%) with NLST participants (52.0%).<sup>4</sup>
- Baseline LDCT LCSs are more likely to discover significant abnormalities and higher rates of late-stage lung cancer.<sup>6</sup>
- The information provided in the article relies on procedure codes, which are known for their inherent inaccuracies. Utilizing coding data poses challenges in accurately reporting follow-up testing and may incorrectly be attributed to screening. Even patients with a negative screen had a 10+% frequency of LDCT and chest CT over the next year which would not be attributable to screening.<sup>7</sup>
- In the PROSPR study, six major complications happened in people without lung cancer, this is only 0.07% of participants. However, most of the complications occurred in people eventually diagnosed with lung cancer.

Easy-to-use evidence-based resources developed by the American College of Radiology and the American Cancer Society, National Lung Cancer Roundtable, such as the Quick Reference Guides on managing [Incidental Findings](#) in LCS and appropriate [Billing & Coding](#) in LCS are now available for LCS programs. Wider distribution and use of these guides would help address some of the issues raised in the article.

## References

1. Lee, Elizabeth, and Ella A. Kazerooni. "Lung Cancer Screening." *Seminars in Respiratory and Critical Care Medicine*. Vol. 43. No. 06. 333 Seventh Avenue, 18th Floor, New York, NY 10001, USA: Thieme Medical Publishers, Inc., 2022.
2. Henschke, Claudia, et al. "Perspective on Management of Low-Dose Computed Tomography Findings on Low-Dose Computed Tomography Examinations for Lung Cancer Screening. From the International Association for the Study of Lung Cancer Early Detection and Screening Committee." *Journal of Thoracic Oncology* (2023).
3. American College of Radiology Committee on Lung-RADS. Lung-RADS Assessment Categories 2022. Available at <https://www.acr.org/-/media/ACR/Files/RADS/Lung-RADS/Lung-RADS-2022.pdf>. Accessed on Jan. 19, 2023.
4. National Lung Screening Trial Research Team. "The National Lung Screening Trial: Overview and Study Design." *Radiology* 258.1 (2011): 243–253.
5. Kaminetzky, Mark, et al. "Effectiveness of Lung-RADS in reducing false-positive results in a diverse, underserved, urban lung cancer screening cohort." *Journal of the American College of Radiology* 16.4 (2019): 419-426.
6. Finigan, James H., and Jeffrey A. Kern. "Lung Cancer Screening: Past, Present and Future." *Clinics in Chest Medicine* 34.3 (2013): 365–371.
7. Hoffman, Richard M., et al. "Lung Cancer Screening With Low-Dose CT: A Meta-Analysis." *Journal of General Internal Medicine* 35 (2020): 3015–3025.