

## RATIONALE:

Lung cancer screening (LCS) with low-dose CT (LDCT) reduces disease mortality,<sup>1,2</sup> but real-world implementation is limited by LDCT access and workflow impediments.<sup>3</sup> We integrated DELFI FirstLook Lung (FLL),<sup>4\*</sup> a blood-based genomic lung screening test, into our electronic health record (EHR) and clinical workflow to improve LCS rates in our health system, focusing on patients who were behind on screening.

## METHODS:

From Oct 2024 through Nov 2025 we offered FLL to USPSTF-eligible patients. EHR-based decision support identified eligible patients, prompted providers and included patient navigation. Our payor partner provided reimbursement for FLL (with potential co-pay).

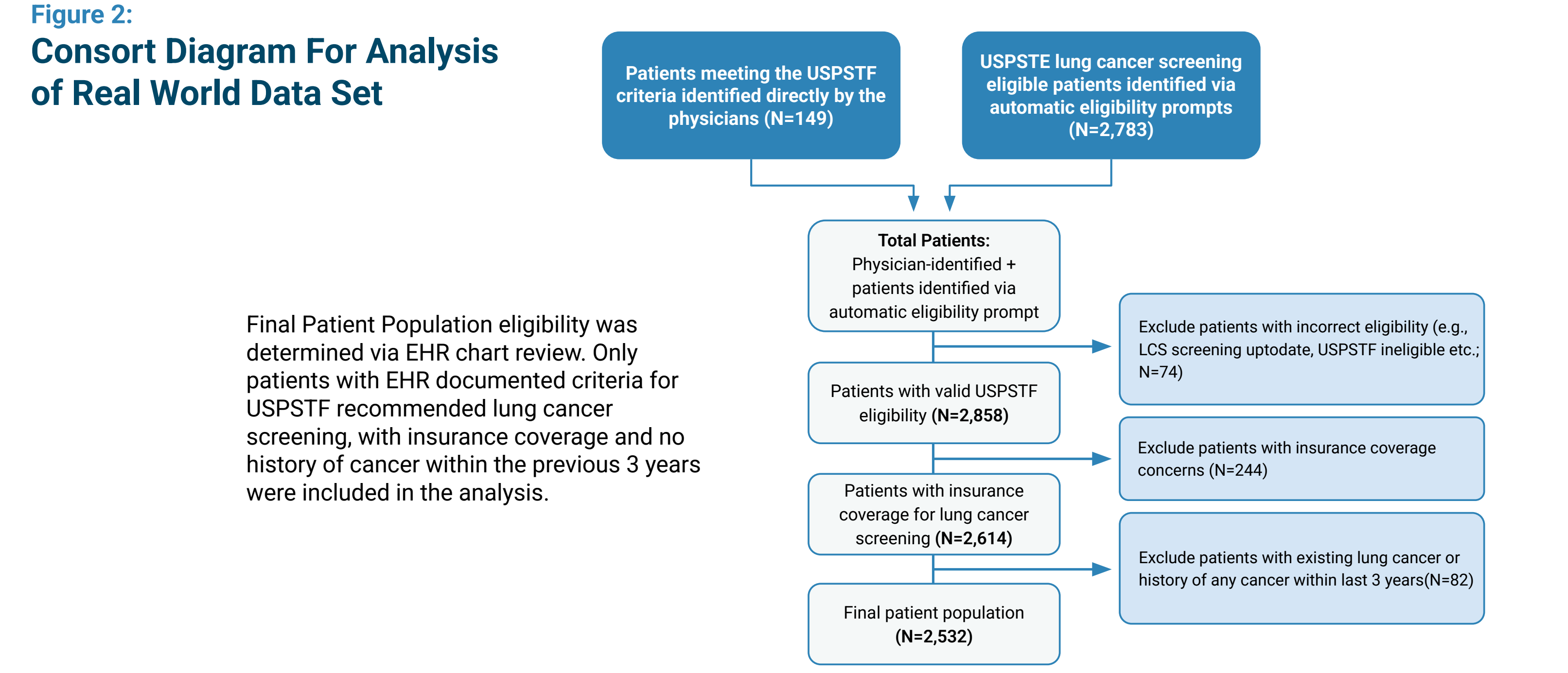
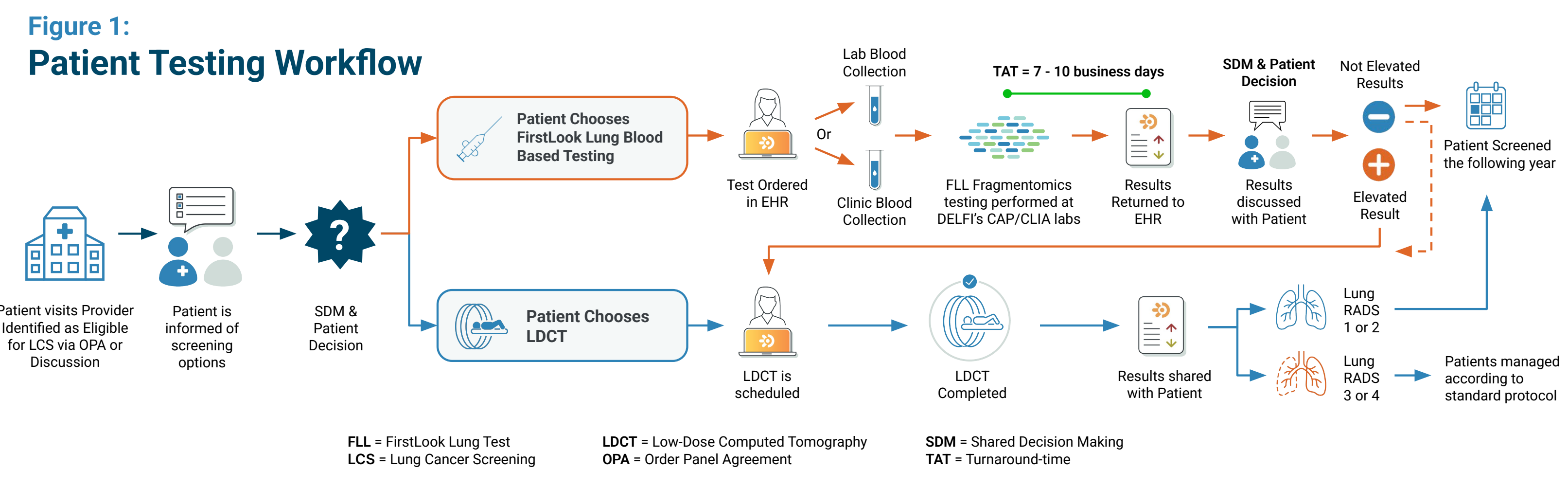
Patient demographics were determined by EHR chart review. Analysis of socioeconomic status was performed by mapping patients zip codes to the CDC/ATSDR Social Vulnerability Index (SVI)<sup>5</sup> which assesses vulnerability across four domains that are relevant to lung cancer screening access: socioeconomic status; household characteristics; racial and ethnic minority status; housing type and transportation.

## RESULTS

- Of the 2,532 patients eligible for LCS (1,331 of whom were LCS naïve), 1,228 (48.5%) were screened either with FLL (n=335) or primary LDCT (n=893) over the 14-month intervention, representing a approximately 2-fold increase compared to the prior year screening rate of 24.1%, and substantially exceeding the historical rate of increase of +3.4 pp per year.
- FLL testing reached a higher proportion of screening-naïve patients than did primary LDCT (63.9% vs. 31.5%, p<0.001). 90% of FLL-screened patients were overdue on their testing and/or annual repeat at the time of testing.
- When compared to patients who were offered primary LDCT screening, having the FLL blood test was associated with greater LDCT completion rates (73.6% vs. 58.6%, p<0.001) and more rapid follow through to LDCT completion both at beginning and end of period (51 down to 15 days and 95 down to 35 days, respectively).
- Suspicious findings on LDCT were more common following FLL Elevated results than primary LDCT (RADS-3: 7.6% vs. 6.6%, RADS-4 or higher: 10.4% vs. 6.1%).

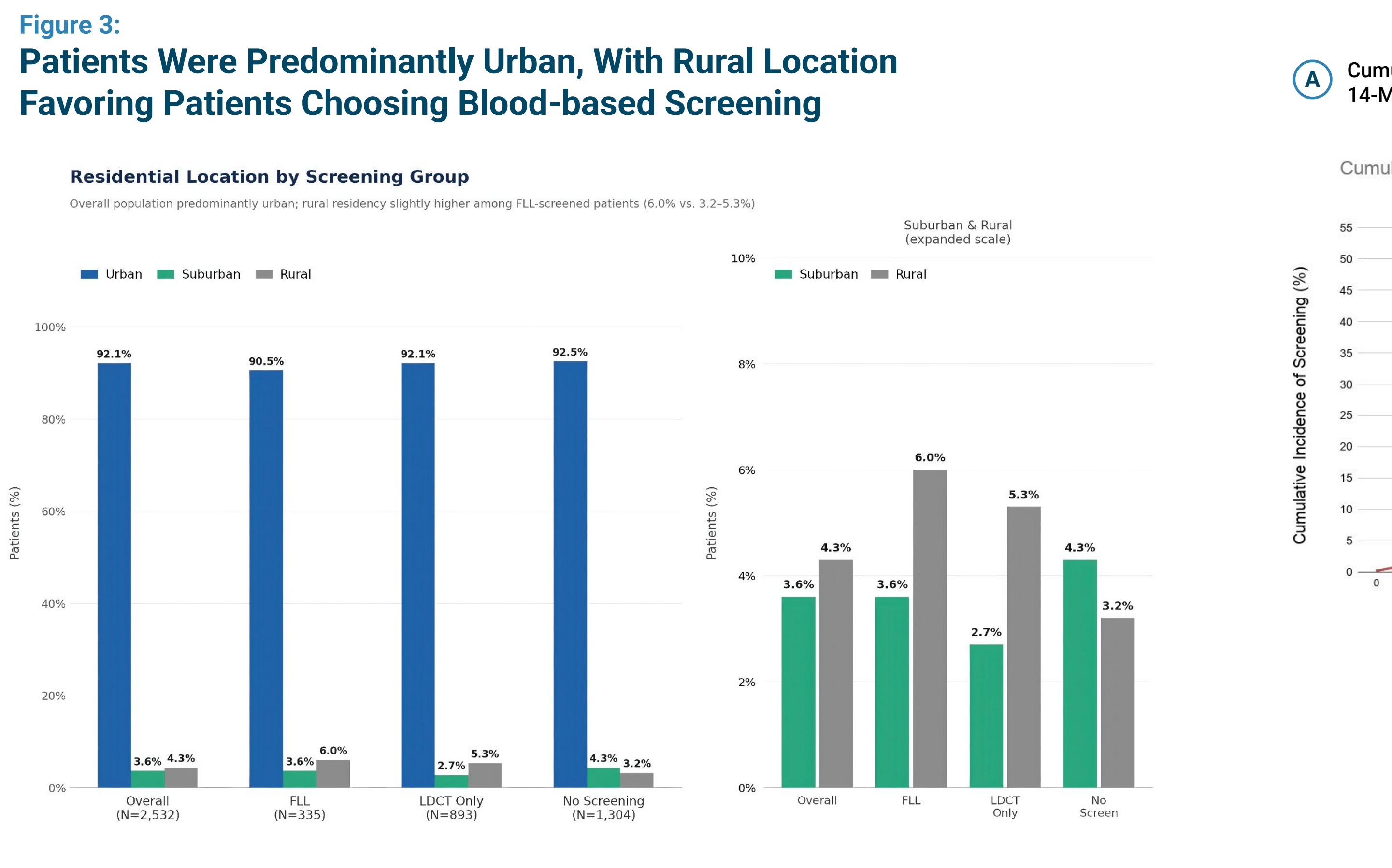
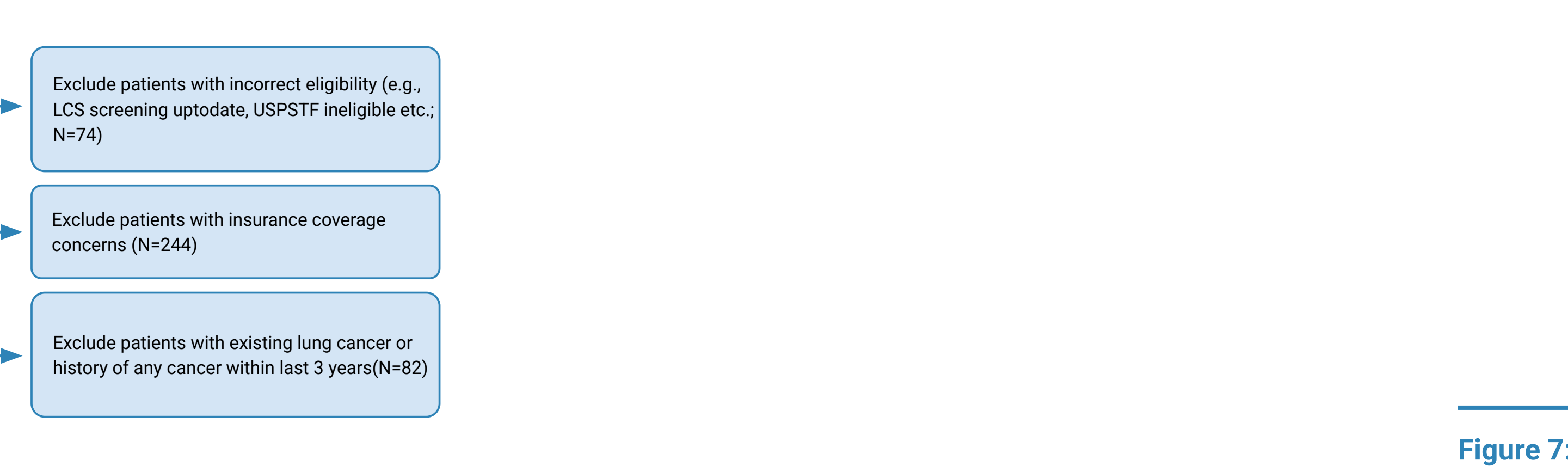
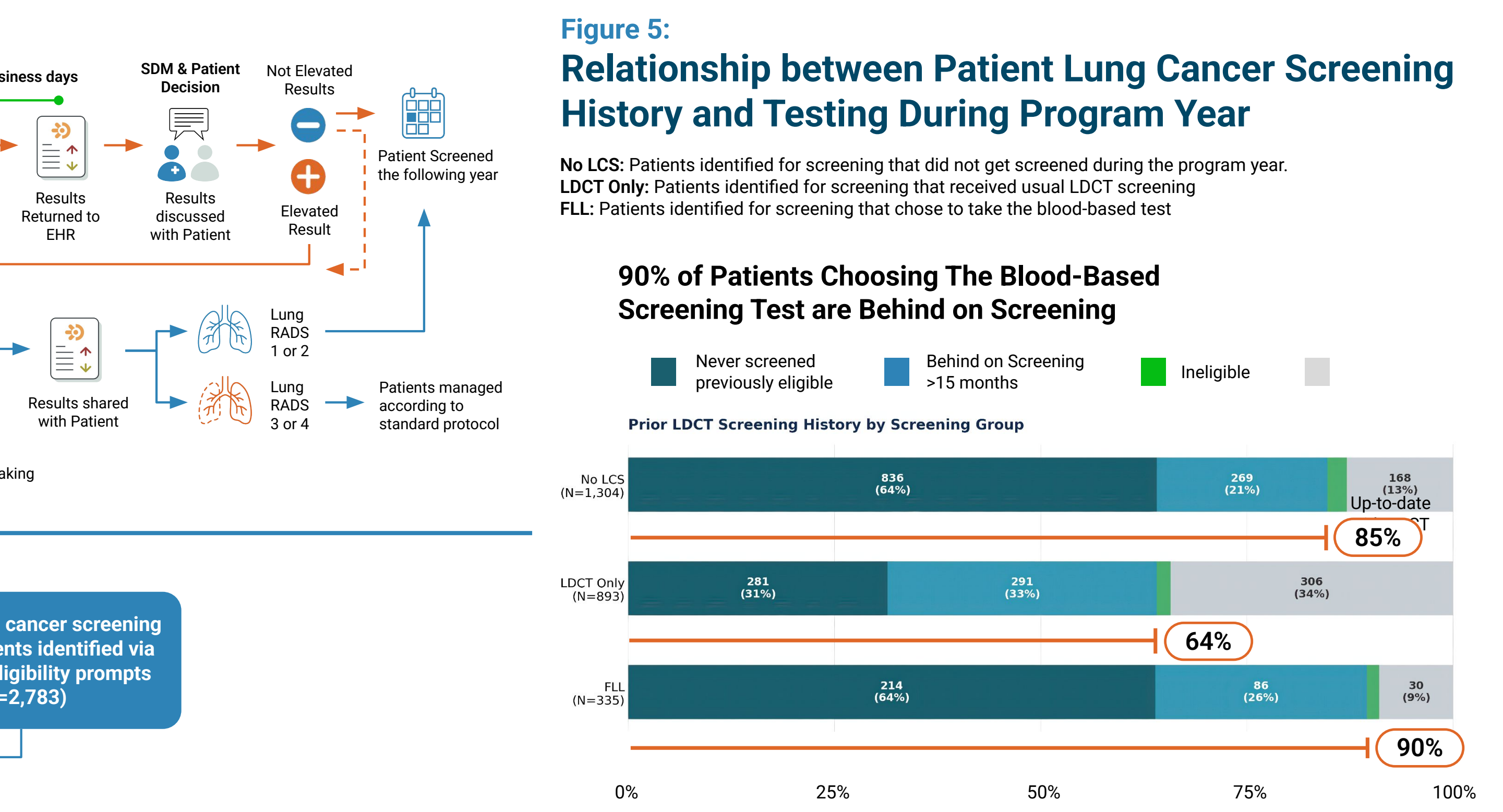
**REFERENCES:**  
 1: "Reduced Lung-Cancer Mortality with Low-Dose Computed Tomographic Screening". The NLSST Research Team. N Engl J Med 2011;365:395-409. VOL. 365 NO. 5  
 2: "Reduced Lung-Cancer Mortality with Volume CT Screening in a Randomized Trial". Harry J. de Koning, M.D. et al. N Engl J Med 2020;382:503-513. VOL. 382 NO. 6  
 3: "Understanding patient barriers and facilitators to uptake of lung screening using low dose computed tomography: a mixed methods scoping review of the current literature" Cavers et al; Respiratory Research volume 23: 374 (2022)  
 4: "Clinical Validation of a Cell-Free DNA Fragmentome Assay for Augmentation of Lung Cancer Early Detection". Mazzone et al Cancer Discov. 2024 Jun 6; 14(11):2224-2242.  
 5: Centers for Disease Control and Prevention/ Agency for Toxic Substances and Disease Registry/ Geospatial Research, Analysis, and Services Program. CDC/ATSDR Social Vulnerability Index 2022 Database United States. https://www.atsdr.cdc.gov/place-health/php/svi/index.html Accessed on April 29, 2026 https://www.atsdr.cdc.gov/placeandhealth/svi/data\_documentation\_download.html Accessed on April 29, 2026.

**\*DISCLAIMER**  
 The FirstLook Lung test is a laboratory-developed test. This test was developed, and its performance characteristics were determined by DELFI Diagnostics. It has not been cleared or approved by the US Food and Drug Administration (FDA). The laboratory is regulated under the Clinical Laboratory Improvement Act (CLIA) as qualified to perform high-complexity clinical tests. The test is used for clinical purposes. It should not be regarded as investigational or for research.

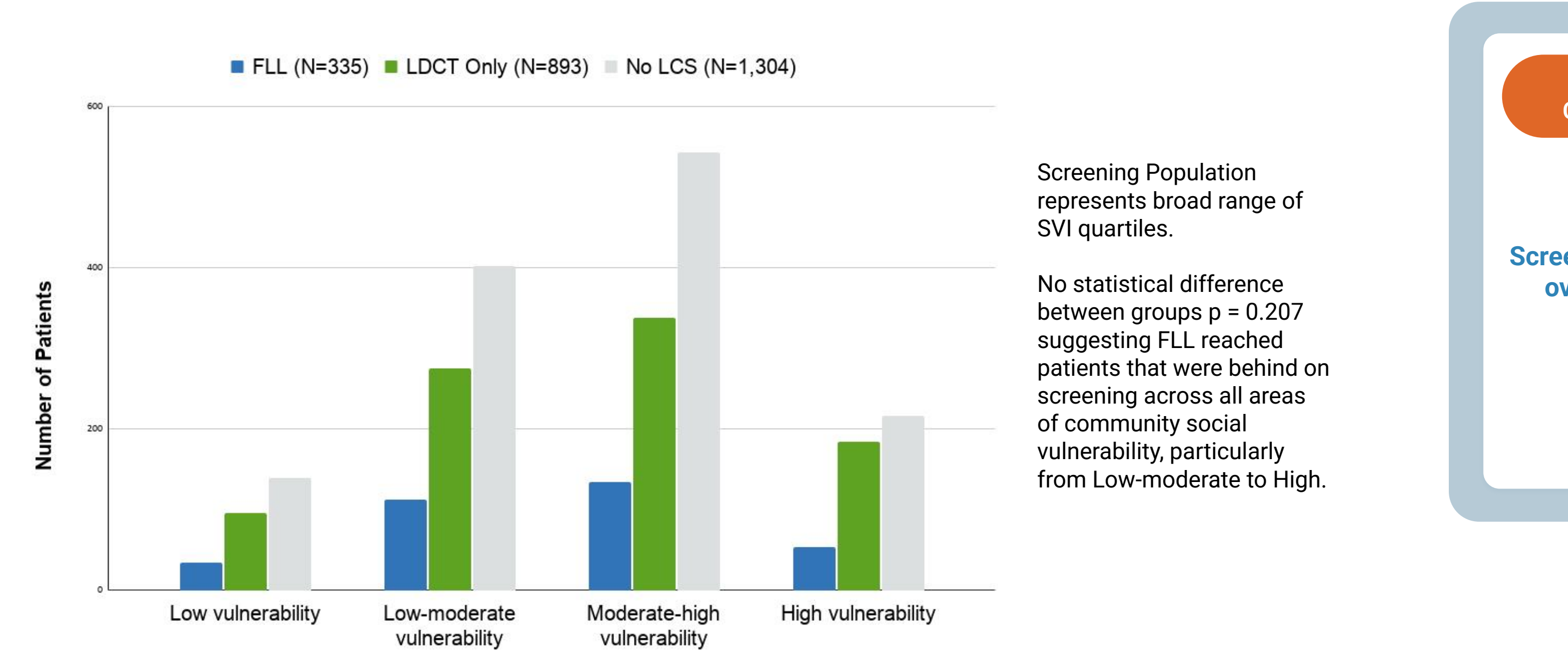


**Table 1: Demographics Across the Eligible Population Within Insurance Coverage Plan Invited for Screening With Results (from Oct 2024 through end Nov 2025; with complete demographic information in EHR)**

	Overall (N=2532)	FLL (N=335)	LDCT Only (N=893)	No LCS (N=1304)	p value
<b>Age</b>	59.6 ± 5.0	60.4 ± 5.9	59.9 ± 4.7	59.2 ± 4.9	<0.001
<b>Sex</b>					
Male	1099 (43.4%)	157 (46.9%)	386 (43.2%)	556 (42.6%)	0.376
Female	1433 (56.6%)	178 (53.1%)	507 (56.8%)	748 (57.4%)	
<b>Race</b>					
White	2340 (92.4%)	321 (95.8%)	819 (91.7%)	1200 (92.0%)	
Black	137 (5.4%)	13 (3.9%)	53 (5.9%)	71 (5.4%)	0.069
Other/Declined	55 (2.2%)	1 (0.3%)	21 (2.4%)	33 (2.5%)	
<b>Ethnicity</b>					
Hispanic or Latino	17 (0.7%)	2 (0.6%)	2 (0.2%)	13 (1.0%)	
Not Hispanic or Latino	2470 (97.6%)	326 (97.3%)	873 (97.8%)	1271 (97.5%)	0.870
Declined	45 (1.8%)	7 (2.1%)	18 (2.0%)	20 (1.5%)	
<b>Smoking Status</b>					
Current	1348 (53.2%)	181 (54.0%)	412 (46.1%)	755 (57.9%)	<0.001
Former	1184 (46.8%)	154 (46.0%)	481 (53.9%)	549 (42.1%)	
<b>Pack years</b>	35.0 ± 15.9	35.5 ± 16.4	36.3 ± 16.4	34.0 ± 15.3	0.003
<b>Residence</b>					
Rural	109 (4.3%)	20 (6.0%)	47 (5.3%)	42 (3.2%)	
Suburban	92 (3.6%)	12 (3.6%)	24 (2.7%)	56 (4.3%)	0.021
Urban	2330 (92.1%)	303 (90.5%)	822 (92.1%)	1205 (92.5%)	

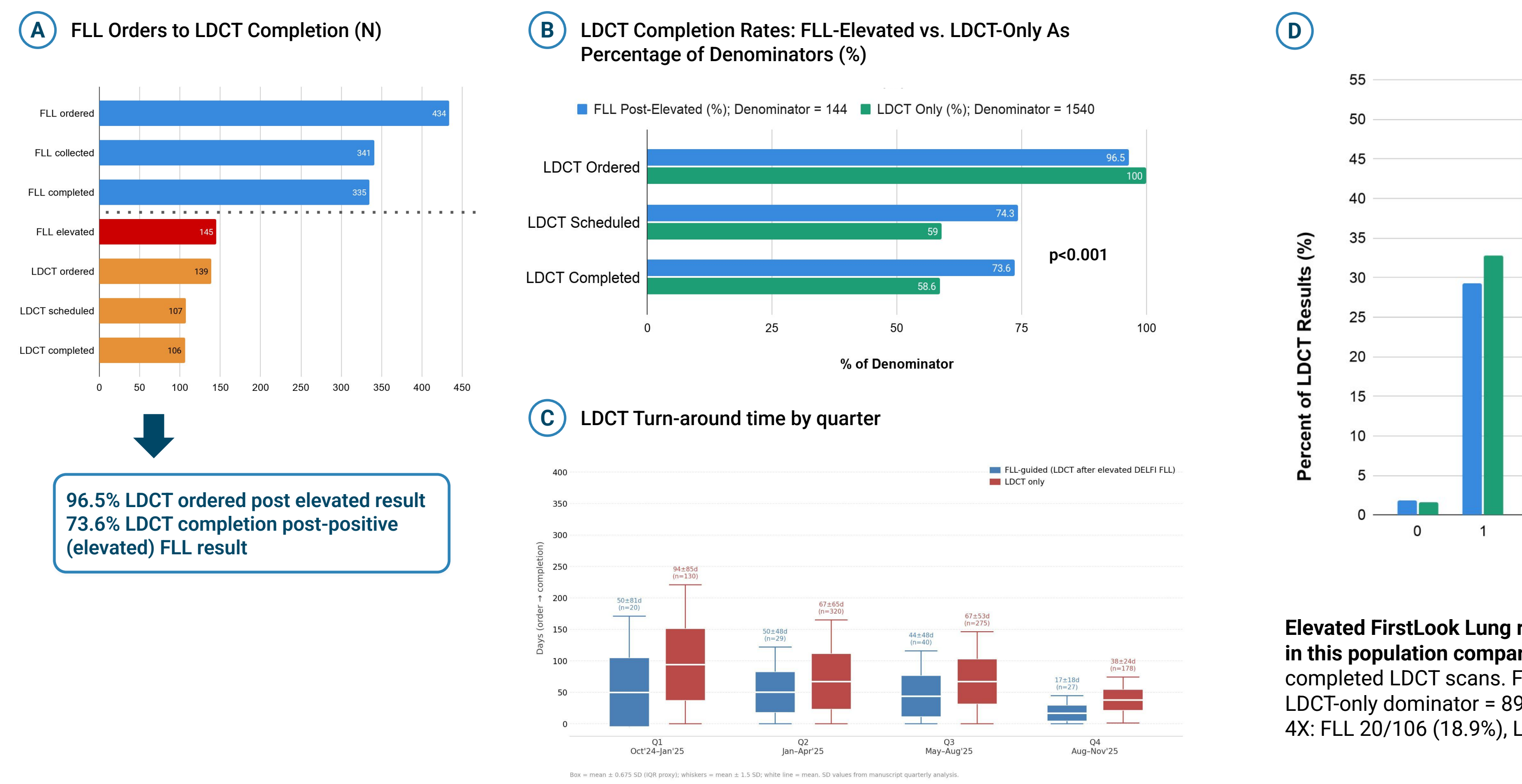


**Figure 4: Screening Modality by Social Vulnerability Index Quartile. Patients Choosing Blood-Based Screening Were from a Broad Spectrum of Community Social Deprivation from Low to High-Vulnerability.**



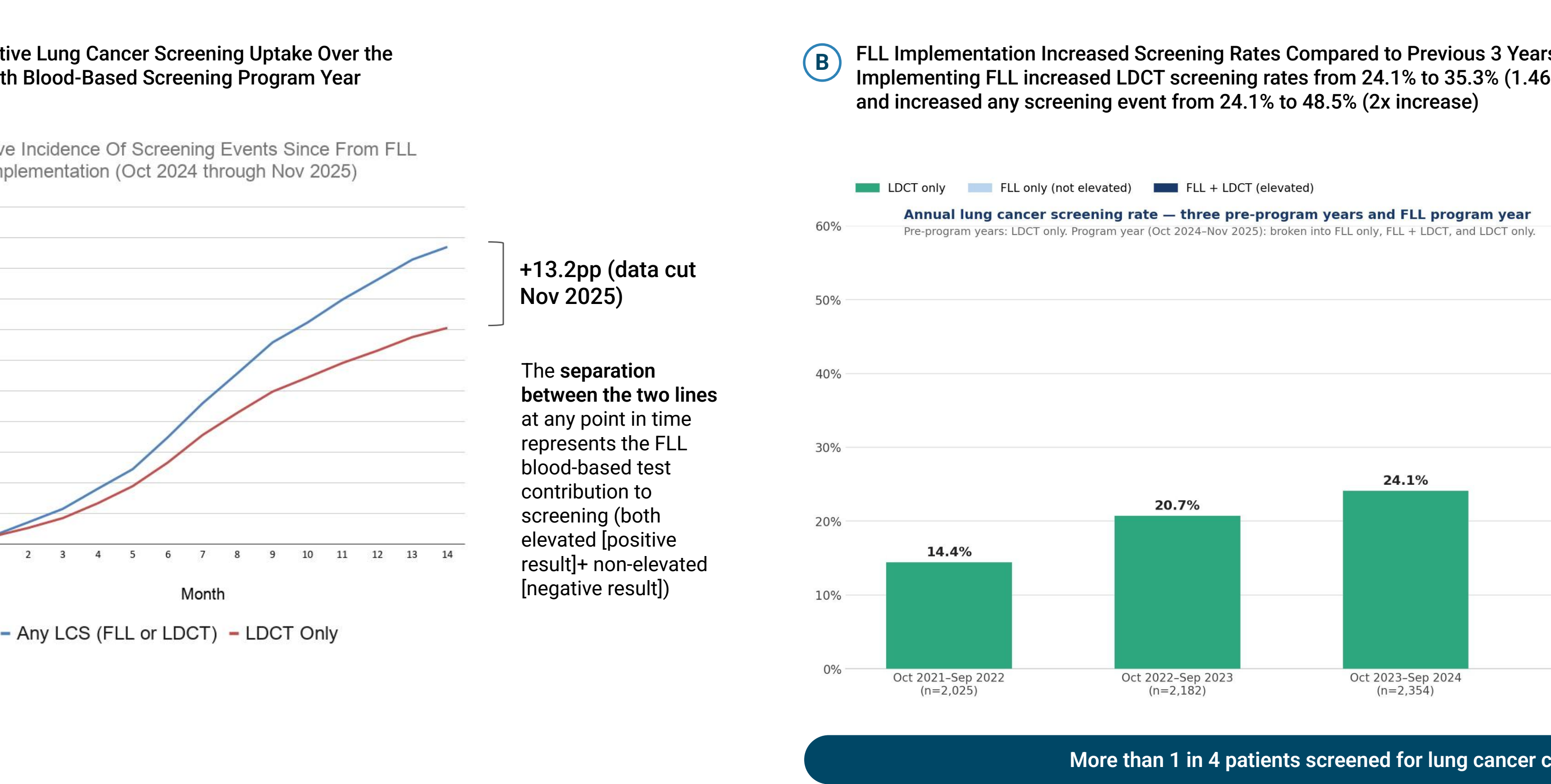
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**Figure 6: Patients Screened by FLL had Greater LDCT Completion Rates, more Rapid Time to Imaging Completion and were associated with more Suspicious LDCT Findings than LDCT Only**



LDCT completion rates (A,B), and time to completion (C) for post-positive FLL results versus LDCT-Only. Compared to patients who were offered primary LDCT screening, FLL blood test results of Elevated ("positive") was associated with greater LDCT completion rates (75.4% vs. 57.2%, p<0.001) and faster turnaround time (C).

**Figure 7: Impact of Implementation of a Blood-Based Lung Cancer Screening Program on Real-World Lung Cancer Screening Rates. Screening Rates are Assessed in the Same Population Across 4 Years (adjusted based on USPSTF eligibility by year)**



13.2% of the eligible population chose and completed blood-based Lung Cancer Screening. Of those screened during the program (14 months), 27.5% were screened via FLL. TOTAL LDCT imaging increased +11.2 pp over the previous year (from 24.1% to 35.3% [31.1% + 4.2%]), compared to +3.4pp in prior years. TOTAL any screening event (LDCT Only + FLL) increased +24.4pp over the previous year. pp = percentage points

Any Lung Cancer Screening

**2X**

Screening Rate Increase over previous year

Any LCS: 48.5% vs. 24.1% (previous year)

LDCT Screening

**1.46X**

LDCT Rate Increase over previous year

LDCT: 35.3% vs. 24.1% (previous year)

Follow-through post Positive

Post-positive result LDCT follow-up **73.6%**

Post-negative result LDCT follow-up **5.0%**

L-RADS 3 and Higher Detected

Blood-Based Test followed by LDCT **18.0%**

LDCT Only **12.6%**

**CONCLUSIONS**

- Implementation of FLL ordering as an option for LCS led to overall increases in screening through both blood and primary LDCT.
- The program improved uptake particularly among those who were screening naïve or behind on annual LCS, suggesting blood-based testing was expanding the population being screened rather than displacing individuals otherwise adhering to LDCT.
- The blood-based test was associated with higher and more rapid rates of follow-through testing than LDCT-Only.

Given the slow adoption of LDCT screening in the USPSTF eligible population, this point of care blood-based screening test has the potential to improve uptake and therefore lower lung cancer mortality through higher rates of early detection.