





Occupational lung cancer screening: A Collegium Ramazzini statement

Steven Markowitz MD, DrPH¹  | Knut Ringen DrPH, MHA, MPH²  |
 John M. Dement PhD³ | Kurt Straif MD, PhD, MPH^{4,5} |
 L. Christine Oliver MD, MPH, MS⁶  | Eduardo Algranti MD, MSc, PhD⁷ |
 Dennis Nowak MD⁸ | Rodney Ehrlich MBChB, DOH, FCPHM(SA)(Occ Med), PhD⁹  |
 Melissa A. McDiarmid MD, MPH, DABT¹⁰ | Albert Miller MD^{1,11} | Collegium Ramazzini

¹Barry Commoner Center for Health & the Environment, Queens College, City University of New York, New York, New York State, USA

²CPWR—The Center for Construction Research and Training, Silver Spring, Maryland, USA

³Duke University School of Medicine, Division of Occupational & Environmental Medicine, Durham, North Carolina, USA

⁴ISGlobal, Barcelona, Spain

⁵Boston College, Chestnut Hill, Massachusetts, USA

⁶Dalla Lana School of Public Health, Division of Occupational and Environmental Health, University of Toronto, Toronto, Ontario, Canada

⁷Division of Applied Research, FUNDACENTRO, Sao Paulo, Brazil

⁸Institute and Clinic for Occupational, Social and Environmental Medicine, LMU Klinikum, LMU Munich, CPC Munich, Comprehensive Pneumology Center Munich, #DZL, Deutsches Zentrum für Lungenforschung, Munich, Germany

⁹Division of occupational Medicine, School of Public Health, University of Cape Town, Cape Town, South Africa

¹⁰Division of Occupational & Environmental Medicine, University of Maryland School of Medicine, USA

¹¹Department of Medicine, Mount Sinai School of Medicine, New York, New York State, USA

Correspondence

Steven Markowitz, MD, DrPH, Barry Commoner Center for Health & the Environment, Queens College, City University of New York, New York, NY, USA.

Email: smarkowitz@qc.cuny.edu

KEYWORDS

low dose CT scan, lung cancer screening, occupational lung cancer

1 | SUMMARY

Lung cancer is the most common cause of death from cancer in the world. It is also the most common lethal work-related cancer. After tobacco smoking, occupational exposures present the most frequent specific cause of lung cancer that is amenable to intervention.

Early detection and treatment can identify and cure primary lung cancer. Randomized controlled trials have demonstrated the efficacy of low dose computed tomography (LDCT) screening among persons at high risk of lung cancer. Guidelines for determining eligibility for LDCT screening have been established for the general population but

have largely neglected those for whom occupational exposure to lung carcinogens is a risk factor.

The Collegium recommends that persons at risk for lung cancer from occupational exposures be offered annual LDCT if their cumulative risk of lung cancer approximates the level of risk endorsed by the guidelines promulgated by the United States Preventive Services Task Force (USPSTF) in 2021 and the National Comprehensive Cancer Network (NCCN) in the United States in 2021. At present, these agencies recommend screening for people aged 50 and over who have smoked at least 20 pack-years of cigarettes. The Collegium recommends that additional lung cancer risk factors, including exposure to known or suspected occupational and environmental

This is an open access article under the terms of the [Creative Commons Attribution-NonCommercial-NoDerivs](https://creativecommons.org/licenses/by-nc-nd/4.0/) License, which permits use and distribution in any medium, provided the original work is properly cited, the use is non-commercial and no modifications or adaptations are made.

© 2024 The Authors. *American Journal of Industrial Medicine* published by Wiley Periodicals LLC.

lung carcinogens; family history of lung cancer (especially among first degree relatives and relatives <60 years of age); a personal history of chronic obstructive lung disease, pneumoconiosis, or pulmonary fibrosis; or a personal history of cancer (excluding skin cancer) be considered as part of the risk assessment for eligibility determination for lung cancer screening. Latency, or the period of time since initial occupational exposure (e.g., >15 years) is another factor that should be considered. If the presence of these additional risk factors, in combination with age and smoking history, is associated with a level of risk that meets or exceeds the level of risk identified by the USPSTF and NCCN, then an annual low dose chest CT for lung cancer screening should be offered. We do not favor a specific age cut-off at which to end screening, but we recognize that only persons who are sufficiently healthy and have sufficient life expectancy to undergo diagnostic work-up and potentially curative treatment should be offered screening for lung cancer. In view of the rising risk of occupational lung cancer over time and the potential or actual interaction between occupational lung carcinogens and cigarette smoking even after quitting, screening programs may choose to screen workers with occupational lung cancer risk for prolonged periods after they have quit smoking cigarettes. The Collegium acknowledges that there are uncertainties and assumptions entailed in this approach and that risk assessment for individual workers necessitates application of significant professional judgement. We encourage the implementation of well-organized screening programs that can further our knowledge about optimal occupation-inclusive lung cancer screening strategies.

Workers with a history of exposure to known or suspected lung carcinogens or working in occupations/trades or work tasks that are known to elevate the risk for lung cancer form the target population for lung cancer screening. Important examples of lung carcinogens include asbestos, silica, diesel exhaust, welding fumes, selected metals, and radiation.

There are well established, evidence-based procedures for the performance of lung cancer screening, preferably in well-organized programs, that apply recognized criteria for cancer screening:

- Screening participants should be provided with complete and comprehensible information about risks and benefits.
- Screening should be offered annually and continuously.
- Screening should be achieved through the application of low dose computed tomography (LDCT) to minimize the radiation dose delivered.
- Proper CT scan interpretation should be performed by experienced radiologists or other well-trained readers.
- Prompt, appropriate follow-up of abnormal CT scans involving relevant medical expertise is mandatory.
- Patients who are current smokers should be offered smoking cessation programs.

The Collegium calls upon occupational health and medical professionals and stakeholders (governments, employers, insurance companies, and labor unions) to identify worker populations that have excess lung cancer risk, to promote lung cancer screening, and

to develop and support well-organized programs to conduct such screening in these populations.

While elimination or minimization of exposure to lung carcinogens in the workplace through environmental controls is critical for lung cancer prevention, lung cancer screening is an essential secondary intervention for reducing deaths and disabling disease from exposure to workplace lung carcinogens.

2 | EVIDENCE-BASIS FOR THIS STATEMENT

2.1 | Burden of occupational lung cancer

Lung cancer is the most common cause of death from cancer in the world, causing one in five (20.4%) cancer deaths in 2019.¹ It is the most common cause of cancer death for males in most countries, including low-, middle-, and high-income nations, and the most frequent cause of cancer death among women in China, the United States, Australia, Scandinavia, and Canada. Tobacco smoking is the dominant cause of lung cancer, and the maturity of the cigarette smoking epidemic and variable uptake and adoption of smoking cessation determines much of the geographic and gender variation in lung cancer incidence and mortality.²

Lung cancer is also the dominant cause of occupational cancer (excluding nonmelanoma skin cancers), causing more than 50% of all workplace-related cancers.³ A recent analysis associated with the Global Burden of Disease Study 2016 estimated that 300,000 lung cancer deaths occurred as a result of exposure to 10 IARC Group 1 lung carcinogens in 2016, representing 86% of all occupational cancer deaths.⁴ Work-related lung cancer deaths increased 55% from 1990 to 2016, from an estimated 193,000–300,000 deaths per year (GBD 2016 Occupational Carcinogens Collaborators 2020). Excellent reviews of occupational cancer in general are readily available.^{5–7}

Occupational lung cancer remains grossly neglected by public health surveillance, clinical medicine, and worker compensation systems, despite its enormous burden of illness and death. Studies in diverse populations and industries across three continents (Asia, Europe, and North America) have demonstrated that a very small fraction—less than 3%—of the total number of estimated occupational lung cancers have been attributed to occupation. In Korea, where an estimated 630 to 1181 occupational lung cancers occur annually, only 179 work-related lung cancers, or 10 per year on average, were compensated by the Korean national worker compensation system over a nearly two-decade period.^{8,9} In Great Britain, where 5442 occupational lung cancer cases are estimated to occur each year,¹⁰ only 21 cases per year (or 392 cases over a 19 year period, 1996–2014) were recorded in Surveillance of Work-Related and Occupational Respiratory Disease (SWORD), a national voluntary reporting system.¹¹ Similarly, in Canada, of the estimated 4150 annual occupational lung cancer cases, only 120 occupational lung cancers were compensated each year between 2005 and 2009.^{7,12}

2.2 | Exposure to occupational lung carcinogens

Over the past five decades, the International Agency for Research on Cancer (IARC) has identified 20 IARC Group 1 occupational lung carcinogens (substances or mixtures) and an additional 7 occupations, industries or work processes in which occupational epidemiology studies were instrumental in establishing specific lung carcinogenicity.⁵ These agents, occupations and industries are listed in Table 1, adapted from IARC sources.^{5,13} Four in ten of all agent-specific IARC Group 1 carcinogens cause lung cancer. In addition, two-thirds of all occupations, industries, or processes that cause occupational cancer cause lung cancer (Table 1).

Further, there is limited evidence for an association with lung cancer of numerous other exposures, though less broadly recognized within the occupational health community. They include cobalt,^{2,3,7,8} tetrachlorordibenzo-*para*-dioxin (dioxin), and high temperature frying emissions and total eight agents or mixtures and four occupations, industries or processes (Table 1).¹³

The number of occupational lung carcinogens are increasing. In the past decade alone, IARC has added common exposures such as diesel engine exhaust (2013), outdoor air pollution (2016), and welding fumes (2017) to its Group 1 list of carcinogens (Table 1).^{5,13} For

additional carcinogens, there is limited evidence for an association with lung cancer: emissions from combustion of biomass fuel (2010); bitumens from roofing (2013); diazinon (2017); and hydrazine (2018).

The occupational lung cancer burden is likely to grow. Only a small fraction of the tens of thousands of chemical agents in commercial use have been evaluated for toxicity. In five decades, IARC has evaluated more than 1000 agents, occupations and industries, but found that available scientific studies are inadequate or lacking in approximately one-half of the evaluations.^{5,14} For context, there are an estimated 86,000 chemicals in the United States Environmental Protection Agency's Toxic Substances Control Act Inventory.¹⁵ Given the frequency of exposure of the respiratory system to inhaled toxicants and the demonstrated carcinogenicity of many chemical agents, it is likely that only a fraction of occupational lung carcinogens has been identified and the total burden of occupational lung cancer remains undefined.

Exposure to occupational lung carcinogens has been and remains reasonably common. National and cross-national surveys of workplace exposures have been conducted in high income countries for 4 decades, including the US National Occupational Hazard and Exposure Surveys (1972–1974 and 1981–1983); CAREX (carcinogen exposure) project in the European Union (1990–1993)¹⁶; FINJEM (Finnish job-exposure matrix) system in Finland¹⁷; and the Canadian version of FINJEM.¹⁸

TABLE 1 IARC agents and processes with sufficient and limited evidence for lung cancer causation.

	Sufficient	Limited
Agent	Arsenic and inorganic arsenic compounds Asbestos (all forms) Beryllium and beryllium compounds Bis(chloromethyl)ether; chloromethyl methyl ether (technical grade) Cadmium and cadmium compounds Chromium (VI) compounds Coal, indoor emissions from household combustion Coal tar pitch Engine exhaust, diesel Nickel compounds Outdoor air pollution Particulate matter in outdoor air pollution Plutonium Radon-222 and its decay products Silica dust, crystalline, in the form of quartz or cristobalite Soot Tobacco smoke, secondhand Welding fumes X-, and Gamma-radiation	Acid mists, strong organic Benzene Biomass fuel (primarily wood), indoor emissions from household combustion of Bitumens, occupational exposure to hard bitumens and their emissions during mastic asphalt work alpha-Chlorinated toluenes (benzyl chloride, benzotrchloride, benzyl chloride) and benzoyl chloride (combined exposures) Cobalt metal with tungsten carbide Creosotes Diazinon Hydrazine Nonarsenical insecticides (occupational exposures in spraying and application of) Silicon carbide, fibrous 2,3,7,8 Tetrachlorordibenzo- <i>para</i> -dioxin Trivalent antimony Uranium, mixture of isotopes
Occupation, industry or process	Acheson process, occupational exposures associated with Aluminum production Coal gasification Coke production Hematite mining (underground) Iron and steel founding Painter (occupational exposure) Rubber manufacturing industry	Art glass, glass containers and pressed ware (manufacture of) Carbon electrode manufacture Frying, emissions from high-temperature Printing processes (occupational exposures in)

Source: IARC: World Health Organization website.¹³ Adapted from Markowitz and Dickens.²¹

The most prevalent occupational lung carcinogens in high income countries over the past 30 years have been diesel exhaust, welding fumes, and silica. Based on data from Europe, Finland, and Canada, more than 2% of the employed population has been exposed to each of these three mixtures or agents. This proportion has not changed in the past three decades. Exposure to asbestos had been a dominant exposure in these countries, but its use declined markedly in recent decades due to widely accepted bans and restrictions. Asbestos exposure continues in these countries, however, due to large quantities of asbestos-containing materials still in place. For middle- and low-income countries, national estimates of the prevalence of exposure to occupational lung carcinogens have not been identified. Given the extent and lack of adequate regulation of manufacturing, mining, and construction, exposures to said agents is likely to be more common and at higher levels than in high income countries.

For the purpose of lung cancer screening, workplace exposures that were prominent 20–40 years ago are highly relevant today due to the latency of asbestos-related lung cancer. Asbestos exposure was common in worksites in many high-income countries before the 1980s, though exposure in recent decades has declined. The prior and continuing high use of asbestos in China, Russia, India, and selected other countries is almost certainly associated with elevated risk of asbestos-related lung cancer for large populations of workers at, both at present and well into the future.^{19,20} Other highly relevant exposures, such as silica, diesel exhaust, and welding fumes, were

prevalent in the past and remain prevalent in countries of all national income levels.

Salient industries and examples of occupations with current exposure to occupational lung carcinogens are provided in Table 2.²¹ Many construction workers are exposed to the most common lung carcinogens: asbestos, diesel exhaust, silica, and welding. Diesel engine exhaust exposure is highly prevalent among workers who drive or maintain diesel vehicles, including buses, trucks, trains, ships, and heavy equipment. Many workers in mining are exposed to diesel exhaust from mining equipment. Miners and workers in many manufacturing industries continue to have exposure to carcinogenic metals and silica.

2.3 | Smoking and chronic lung disease amplify occupational lung cancer risk

2.3.1 | Cigarette smoking

Occupation and, more generally, social class, are closely associated with cigarette smoking. In the United States, one-quarter or more of workers in construction, manufacturing, mining, and transportation smoke cigarettes compared to 10% of workers in professional or managerial positions.²² In China, for example, the prevalence of smoking among male machine operators (67%) was nearly twice that of male medical/health personnel or teaching staff (36%–38%).²³

2.3.2 | Interaction of smoking and occupational lung carcinogens

It is well-established that occupational lung carcinogens and cigarette smoke act in concert in some circumstances to increase the risk of lung cancer. They share mechanistic pathways and have been repeatedly shown in epidemiologic studies to increase lung cancer risk above that expected by the presence of each risk factor alone. Asbestos is the best-known example of this phenomenon. Asbestos frequently shows at least a supra-additive interaction with smoking in determining lung cancer risk.^{24–28} Several large studies addressing the lung cancer risk among silica-exposed workers have been completed in the past decade, generally suggesting a supra-additive effect with cigarette smoke.^{29–33} Other occupational lung carcinogens that have been studied for interaction include diesel exhaust^{34,35}; and radon.^{36–38}

2.3.3 | Chronic lung disease and lung cancer risk

Occupational exposures also indirectly increase lung cancer risk by causing chronic lung diseases, namely, chronic obstructive lung disease (COPD) and pneumoconioses, such as silicosis and asbestosis.^{21,39} In fact, since occupational exposures to vapors, gases, dusts, or fumes raise the risk of COPD,⁴⁰ for example, the occupational contribution to lung cancer should be considered more broadly than simply the role of the occupational agents causing lung cancer.

TABLE 2 Common occupational lung carcinogens by industry and selected occupations.^a

Industry	Lung carcinogen	Examples of occupation
Manufacturing	Silica, chromium, nickel, cadmium	Metal fabricators, assemblers Metal processors, shaping workers Clay, stone, glass processors Forging workers Boilermakers, platers
Construction	Silica, diesel exhaust, painting, welding, coal-tar pitch, asbestos Outdoor air pollution	Excavators Welders Painters Plumbers Other construction
Transportation	Diesel exhaust, PAH, outdoor air pollution	Bus drivers Truck drivers Mechanical maintenance
Mining, oil, gas extraction	Silica, diesel exhaust radon	Drillers, blasters Miners, quarry workers Mineral ore treaters

^aOccupations listed are examples of workers with exposure within designated industries and do not represent a complete list of such occupations.

Source: Adapted from Markowitz and Dickens.²¹

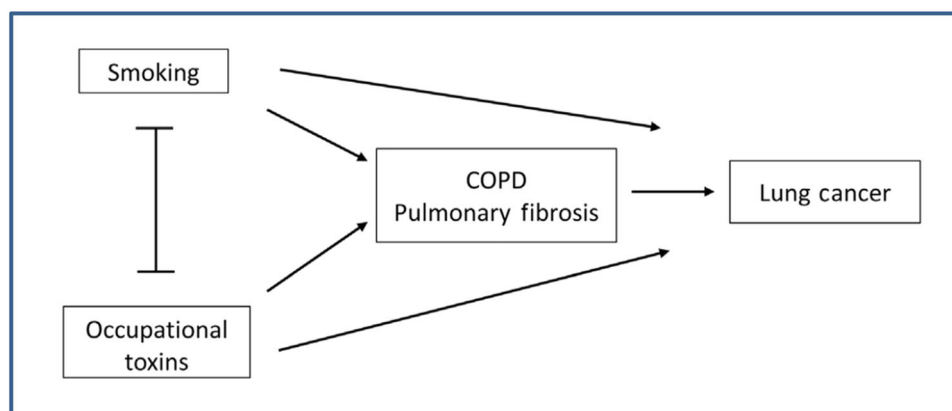


FIGURE 1 Occupation and smoking: nexus of chronic lung disease and lung cancer. Previously published in Markowitz and Dickens.²¹

Smoking works similarly as a major cause of COPD and as an established risk factor for idiopathic pulmonary fibrosis.^{41,42} Figure 1 illustrates the complexity of these relationships. Key aspects of these relationships have been well-studied (e.g., the smoking and asbestos interaction noted above and the contribution of asbestosis and silicosis to risk of lung cancer). Other relationships, such as the interaction between work-related COPD and lung cancer, have received relatively less attention.

3 | LUNG CANCER SCREENING RECOMMENDATIONS IN THE GENERAL POPULATION

3.1 | Low dose CT (LDCT) scan screening studies

Three decades of research provide strong evidence that periodic low dose chest CT scans can identify lung cancers at an early stage and can reduce lung cancer mortality. In Japan and the United States, Sone et al.⁴³ and Henschke et al.⁴⁴ separately demonstrated that low dose chest CT scanning in high risk populations detected ~85% of lung cancers at Stage I. In 2006, Henschke and colleagues further showed that treated early CT-detected lung cancers had excellent survival: a group of 412 Stage 1 lung cancers detected by CT screening had an estimated 88% 10-year survival.⁴⁵ Of those who underwent surgical resection within 1 month of diagnosis, 10-year survival was 92%, 95% CI 88–95%. These remarkable results of nonrandomized studies stimulated intense interest and the initiation of randomized controlled trials of the impact of low dose CT scans on lung cancer mortality.

Two large complementary randomized clinical trials of populations at high risk of lung cancer—the US National Lung Screening Trial (NLST) and the Dutch–Belgian Netherlands–Leuven Longkanker Screenings Onderzoek (NELSON)—conclusively demonstrated that periodic low dose chest CT scanning reduces lung cancer mortality.^{46,47}

The NLST, conducted by the United States National Cancer Institute, included 53,454 enrollees aged 55–74 who had smoked at least 30 pack-years and, for former smokers, had quit within the past 15 years. The CT versus chest X-ray (CXR) study arms were screened

annually for 5 years and followed for a median of 6.5 years. The CT scan screening arm showed a 20% reduction in lung cancer mortality versus the CXR screening arm.⁴⁶

The NELSON trial included 13,195 men and 2594 women aged 50–74 years who had smoked at least 15–20 pack-years and, if former smokers, had quit 10 or fewer years before the entry date into the study. NELSON compared an intervention group who underwent four rounds of CT screening (baseline, year 1, year 3, and year 5.5) with a reference group who had no screening; all were followed for at least 10 years. NELSON observed a 24% and 33% lung cancer mortality reduction among men and women, respectively, in the trial.⁴⁷

Neither the NLST nor the NELSON trials were designed to evaluate the efficacy of lung cancer among screenees defined other than by age and smoking. Other factors would include occupational exposures, chronic lung disease, family history of lung cancer, personal history of cancer, or environmental exposure to radon, air pollution, or other nonoccupational toxins.

3.2 | Lung cancer prediction models

An alternative to the use of age and smoking history alone to estimate lung cancer risk and to determine screening eligibility is the employment of a broader set of lung cancer risk factors in lung cancer risk models. More than 20 lung cancer risk prediction models based on large lung cancer data sets (e.g., prostate, lung, colorectal, and ovarian cancer screening trial [PLCO, developed in North America]⁴⁸; liverpool lung project [LLP, developed in England]⁴⁹) have been developed, and many are inclusive of a broader and more detailed set of lung cancer risk factors than the NLST and NELSON trials.⁵⁰ These additional risk factors variably include gender, race, body mass index (BMI), intensity and duration of smoking, number of years since smoking cessation, chronic lung disease, especially chronic obstructive pulmonary disease (COPD), personal history of cancer, family history of lung cancer, education, and asbestos exposure. Among the better-performing models, the only occupational carcinogen included is asbestos, which is part of the Liverpool Lung Project (LLP) and Bach models.^{49,51} No other occupational or environmental exposures have

been included in the risk prediction models. Lung cancer risk calculators derived from these models use no or little information about occupation in determining risk.

In a noteworthy comparison of the risk factor versus risk prediction approaches to the use of LDCT, ten Haaf⁵² applied nine established lung cancer risk prediction models to the large data sets of the NLST and the PLCO and compared results to the risk factor eligibility criteria used in the NLST, using 5- and 6-year lung cancer incidence and mortality as outcomes. The specificity of the risk prediction models versus the NLST criteria were very similar (~62.2%–62.6%), but four models had substantially higher levels of sensitivity (>78%) compared to that of the NLST criteria (71.4%) with respect to lung cancer incidence. The contrast in sensitivity for lung cancer mortality between the NLST criteria and the model predictions was even greater: 73.5% for the NLST versus 85.2% for the PLCO_{m2012} model and 83.8% for the two-stage clonal expansion (TSCE) and Bach models.⁵²

In a US National Cancer Institute study of this issue, investigators compared the lung cancer mortality benefits of a risk prediction-based model versus a USPSTF guidelines-based model (restricted to age and smoking) and concluded that the former approach, which used family history, self-reported emphysema, body mass index, age, and a broader range of smoking history as eligibility criteria, prevented a greater number of lung cancer deaths than a model based on USPSTF screening guidelines (see below).^{53,54}

3.3 | Current lung cancer screening recommendations

3.3.1 | United States

Based principally on the results of the NLST, the United States Preventive Service Task Force (USPSTF) recommended in 2013 that annual low dose CT scanning be offered to individuals at high risk of lung cancer, who were defined as people aged 55–74 years who had smoked at least 30 pack-years of cigarettes and currently smoke or quit less than 15 years previously. In 2021, after publication of the NELSON trial results, the USPSTF revised its recommendations for annual LDCT eligibility to include people aged 50–80 years who have at least a 20 pack-year smoking history and currently smoke or have quit within the past 15 years.⁵⁵ These recommendations were adopted by the US federal government and private insurance companies for use in health insurance coverage and clinical preventive practice.

3.3.2 | Europe

Recommendations based in Europe to date have adopted a more heterogeneous approach. In a March 2022 report prepared by Science Advice for Policy by European Academies (SAPEA), a consortium effort of European Scientific Academies and released, lung cancer screening with LDCT is recommended, using either a combination of age and smoking or the PLCO_{m2012} model.⁵⁶ An

expert group of European physicians and scientists who are leaders in lung cancer screening in Europe have developed a set of consensus recommendations on lung cancer screening, which were published in June 2020.⁵⁷ They recommended the use of lung cancer risk thresholds (e.g., 1.51% over a 6-year period) applied to results of risk prediction models to determine who should be eligible for LDCT-based lung cancer screening. Of the numerous risk prediction models that have been developed, the expert group favored those derived from the Liverpool Lung Project (LLP_{v2}) and the Prostate, Lung, Colorectal, and Ovarian Cancer Screening Trial (PLCO_{m2012}).

3.3.3 | China

In China, a multidisciplinary lung cancer early detection and treatment expert group (appointed by the Chinese National Health and Family Planning Commission) established the China National Lung Cancer Screening Guidelines, recommending annual lung cancer screening with LDCT for people aged 50–74 years who have at least a 20 pack-year smoking history and who currently smoke or have quit within the past 5 years.⁵⁸

The latest Chinese guidelines developed by the National Cancer Center of China recommends lung cancer screening eligibility including: (1) current smokers with ≥30 pack-years or former smokers with ≥30 pack-years who have quit within 15 years; (2) secondhand smokers who have lived or worked with smokers for at least 20 years; (3) people with COPD; (4) participants who have exposure ≥1 year to asbestos, radon, beryllium, chromium, cadmium, nickel, silica, or soot; (5) people with first degree relatives have confirmed lung cancer.⁵⁹

3.3.4 | South Africa

The South African Thoracic Society recently issued recommended guidelines for lung cancer screening. They recommend that annual LDCT should be offered to people between 55 and 74 years of age who are current or former smokers (having quit less than 15 years previously) with a history of ≥30-pack years of smoking. Participants should have no history of lung cancer and be in reasonable health and able and willing to be treated for lung cancer. They note that the high prevalence of tuberculosis in South Africa requires that only lung nodules ≥6 mm need follow-up.⁶⁰

4 | DEARTH OF RECOMMENDATIONS FOR OCCUPATIONAL LUNG CANCER SCREENING

4.1 | Current recommendations

Occupation is generally ignored in current screening recommendations, whether they are based on selected risk factors (age and smoking) or on risk prediction models. Exceptions in the US include guidelines

developed by the National Comprehensive Cancer Network (NCCN) and the American Society of Thoracic Surgeons. Their guidelines include using additional risk factors to determine screening eligibility, that is, family history, history of chronic lung disease and occupational exposures. Specifically, NCCN Group 2 eligibility criteria, first used in 2014, recommended screening people aged ≥ 50 years with a 20 pack-year smoking history if they have an additional risk factor for lung cancer, such as exposure to occupational lung carcinogens, chronic lung disease, or a family history of lung cancer and had an aggregate 6 year risk of lung cancer $\geq 1.3\%$.⁶¹ More recently, NCCN recommended a 20 pack-year screening threshold for everyone aged 50 years and over but occupation and other risk factors can additionally be considered in determining eligibility for screening.⁶² Occupational lung carcinogens named include silica, cadmium, asbestos, arsenic, beryllium, chromium, diesel exhaust, nickel, coal smoke, and soot.

In 2014, a group of international experts in asbestos-related diseases met in Helsinki and recommended that the lung cancer risk level associated with the NLST study population be used as the threshold risk level in organized low dose CT scan programs for screening asbestos-exposed workers for lung cancer.⁶³ The Helsinki recommendation was made before completion of the NELSON clinical trial, numerous modeling studies, and revision of the USPSTF eligibility guidelines in 2021.

In 2017, an expert working group in France made recommendations for the application of LDCT for lung cancer screening for workers who have a history of exposure to Group 1 IARC occupational lung carcinogens, including asbestos.⁶⁴ They conducted a scientific literature review and developed an expert consensus on how to identify the magnitude of lung cancer risks associated with these Group 1 carcinogens, alone and in combination with cigarette smoking, and identified the combinations that equaled or exceeded the lung cancer risk associated with NLST eligibility criteria. They assumed a multiplicative (and not a supra-additive or additive) joint effect between the occupational carcinogenic agent and tobacco in estimating the relative risks. They estimated that the relative risk of lung cancer associated with the NLST study eligibility criteria (≥ 30 pack-year history of cigarette smoking) was 30. Using the target relative risk level of 30, exposure to each of the IARC Group 1 lung carcinogens in combination with 20–29 pack years of smoking met or exceeded the target risk level. Never smokers who were occupationally exposed to lung carcinogens at any level did not reach a relative risk sufficient to justify lung cancer screening with the possible exceptions of plutonium and bis-chloromethyl ether.⁶⁴

Since the publication of this set of French recommendations in 2017, the results of the NELSON trial and modeling studies led the USPSTF to lower the age and smoking levels for eligibility for LDCT-based lung cancer screening (\geq age 50 years and ≥ 20 pack-years). The associated estimated relative risk level of lung cancer in the French analysis would be 20. Accordingly, under these French guidelines, workers with intermediate asbestos exposure ≥ 10 years and workers with ≤ 5 years of high asbestos exposure would be recommended for lung cancer screening even among ever smokers with a smoking history of less than 20 pack-years.

4.2 | Empirical studies of LDCT in occupational populations at risk

Studies of lung cancer screening among occupational populations at high risk of lung cancer are limited to date. Two types of occupational settings have been studied: asbestos-exposed groups and US nuclear weapons workers.

4.2.1 | Asbestos

Nine nonrandomized studies of asbestos-exposed populations with ≥ 150 participants reported results of LDCT screening for lung cancer between 2002 and 2019.^{65–74} Age, smoking history, and asbestos exposure were principal or exclusive eligibility criteria with frequent use of broader age and smoking ranges than those used in NLST or NELSON. These studies were included in a published systematic review and meta-analyses by Ollier et al.⁷⁵ and Maisonneuve et al.⁷³

None of these studies used risk prediction models, NLST, or NELSON eligibility criteria to determine eligibility. Comparison with NLST or the NELSON results is precluded by the heterogeneity of the study populations and the lack of sufficient detail on smoking history and age in the published studies.

Combining the screening yield results of these nine studies yielded 86 lung cancers detected among 5548 ever smokers (1.55%) and 6 lung cancers detected among 1787 never smokers (0.33%).

Parameters of asbestos exposure varied widely among the nine studies by industry, occupation, duration of employment, presence, or absence of pleural plaques, and other variables. One-half of published studies screened people only if they had a history ≥ 15 years of asbestos exposure, while one-half used a history of 1 or 5 years of significant asbestos exposure as a screening criterion. The studies with the highest lung cancer detection yields by LDCT also tended to have study populations with the highest prevalence (43%–89%) of pleural plaques on low-dose CT scans.^{67,72,76} The pleural plaques were generally identified by LDCT.

Only one study reported on lung cancer mortality follow-up of application of LDCT among an asbestos-exposed population. Barbone and colleagues examined 9-year mortality follow-up of a nonrandomized study of 926 asbestos-exposed workers who were enrolled in an asbestos surveillance program in Northeastern Italy, a major shipbuilding area. The group had undergone at least two periodic LDCT scans beginning in 2002.^{68,77} They were mostly men with a mean age of 58 years at study onset and a mean duration of 30 years of exposure to asbestos; one-third never smoked, and median pack-years of cigarettes among smokers was 18.5. The comparison group of 1507 people had more smokers but a lower average level of asbestos exposure and underwent chest-rays (CXR). The LDCT study group showed a lung cancer SMR of 0.55 (95% CI: 0.24–1.09), and the CXR study group had a lung cancer SMR of 2.07 (95% CI: 1.53–2.71) in comparison with regional mortality rates. In a Cox proportional hazard analysis adjusting for age, smoking, and asbestos exposure level, the LDCT study group showed a hazard ratio for lung

cancer of 0.41 (95% CI: 0.17–0.96) versus the CXR study group. Differences in the lung cancer risk factor profile between the study groups and the nonrandom nature of the study likely accounted for some of the observed difference in lung cancer mortality.

Loewen and colleagues used LDCT to study a population with mostly environmental exposure to Libby amphibole asbestos in Libby Montana. The population was aged 50–84 years, had a >20 pack-year history of tobacco use, and had a history of asbestos-related pleural or pulmonary fibrosis on a previous high-resolution CT scan. Seventeen lung cancers were detected: 71% were stage 1 (59%) or stage 2 (11%).⁷⁸

In Germany, the German Statutory Accident Insurance offers annual LDCT screening to those who are ≥55 years of age, have smoked >30 pack-years and have been exposed to asbestos ≥10 years before 1985 or who have parenchymal asbestosis or pleural fibrosis. As of mid-1991, 10,306 of 20,253 people who met these criteria and had physician counseling underwent baseline LDCT screening with 5476 undergoing a second round of screening and 1725 have additional screening rounds. Preliminary data show that 133 people in the program have been detected as having lung cancer.⁷⁹

In Australia, Brims and colleagues applied LDCT for 5 years to 1743 people with asbestos exposure, including one-third from the Wittenoom environment (workers or residents). Non-Wittenoom participants had >3 months of cumulative occupational exposure to asbestos or radiographic evidence of pleural plaques. Median cumulative asbestos exposure was modest (0.7 fiber/cc-years, but 61% had pleural plaques and 35% had parenchymal asbestosis on the baseline LDCT scan. Only 7% of the population were current smokers, and one-third of the population never smoked; median pack-years was 20 for ever smokers. LDCT detected 22 lung cancers, and an additional 4 lung cancers were detected in follow-up (3 were interval cancers). Nineteen of the 22 (86.4%) LDCT-detected cancers were stage I lung cancers. The authors applied the 2021 USPFTF, PLCO₂₀₁₂, the LLP, and a combined PLCO₂₀₁₂/LLP eligibility criteria to the study population and found that no more than 25% of the program participants would have been screened under these various alternative criteria. Consequently, no more than 35% of the 26 lung cancers detected by the program would have been detected by application of these alternative eligibility criteria. Brims and colleagues conclude that existing lung cancer screening criteria do not adequately account for occupational exposures.⁷⁴

4.2.2 | Asbestos-exposed never smokers compared to smokers

Kato and colleagues performed a one-time LDCT screening of 2132 asbestos-exposed workers, principally from shipbuilding, construction, and manufacturing sectors in Japan. Eligibility criteria included: (1) work in asbestos manufacturing for ≥1 year; or (2) work in other asbestos-exposed industries for >10 years; or (3) evidence of pleural plaques on CXR or chest CT scan. In total, 89% of participants had

pleural plaques on CT scan. Among 444 never smokers, 3 lung cancers were detected (0.7%), and 42 lung cancers were found among 1651 smokers (2.5%).⁷²

In the Brims and colleagues study summarized above, 4 lung cancers were detected among 596 (0.7%) never smokers, representing 15% of all lung cancers detected in the program. Details regarding the asbestos exposure or findings of radiographic scarring are not available.⁷⁴

4.3 | US nuclear weapons workers

For two decades, Markowitz, Miller and colleagues have used low-dose chest CT scanning to screen ~14,000 workers who had worked at nuclear weapons facilities in 13 mostly nonmetropolitan US communities with variable exposure to asbestos, radiation, beryllium, nickel, chromium, and other toxins. Eligibility criteria included age (≥50 years), smoking, occupation (production, maintenance, or laboratory worker), and, if present, asbestos-related radiographic parenchymal or pleural fibrosis, and/or a positive beryllium lymphocyte proliferation test. In a 2018 report on 7189 of these workers, all of whom had a smoking history, the proportions with screen-detected lung cancer were 0.83% at baseline and 0.51% on annual scan. Stage distribution at diagnosis was favorable: of 80 detected lung cancers, 59% ($n = 47$) were stage 1, and 10% ($n = 8$) were stage 2. Study strengths included high study compliance; high credibility with the study population through labor union cosponsorship; implementation in community settings; excellent follow-up; and use of a standardized protocol with demonstrated quality.⁸⁰

To delineate the occupational contribution to aggregate lung cancer risk, Markowitz and colleagues compared two subgroups of the study population: Group A met NLST eligibility criteria for age (≥55 years) and smoking history (≥30 pack-years); Group B did not meet these NLST study criteria but met National Comprehensive Care Network (NCCN) Group II criteria (age ≥50 years, 20–29 pack-year smoking history, and occupational risk). Both groups had occupational exposures. Group B had a lower overall lung cancer risk compared to the NLST study based on age and smoking profile but had occupational risk of lung cancer. The lung cancer screening yield of Group A (1.5%, 95% CI: 0.88–2.12%) was similar to that of Group B (1.36%, 95% CI: 0.85–1.87%). Both were statistically similar to the screening yield of the original NLST study (1.0%, 95% CI: 0.88–1.12%). These results indicate that, in the presence of occupational risk, younger people with lesser smoking histories could nonetheless benefit from LDCT screening.⁸⁰

Although the Markowitz and colleagues study did not include mortality follow-up, the shift in diagnosis of earlier stage lung cancers with LDCT screening is consistent with the NLST and NELSON trials that demonstrated favorable mortality reductions.⁸⁰

A second study of LDCT screening among US nuclear weapons workers—construction workers—was completed by Welch and colleagues. The study group had a lower age and smoking threshold than the NLST study criteria (age ≥50 years and smoking ≥20 pack-years),

but they had additional lung cancer risk, as defined by 5 years of work in the construction industry (exposure to asbestos, silica, beryllium, chromium, radiation, or welding), evidence of radiographic asbestosis or pleural plaques, or spirometric evidence of chronic obstructive pulmonary disease. Stage distribution among CT-detected cancers was favorable: 20 of 30 (67%) detected lung cancers were stage 1 (57%) or stage 2 (10%) disease. The lung cancer screening yield at baseline scan was 1.7% (21 of 1260 participants), a result that was similar to the NLST results, despite the fact that less than one-half of participants met the NLST eligibility criteria.⁸¹

Dement and colleagues used this nuclear weapons construction worker cohort and a related larger construction worker population to develop a lung cancer risk prediction model (BTMed model) that includes age, gender, race, smoking history, spirometry, chest X-ray finding of parenchymal fibrosis and/or pleural plaques, occupational history of ≥ 5 years of work in construction, body mass index, respiratory symptoms, and personal history of cancer.⁸² Applying the lung cancer screening criteria described above in the Welch and colleagues study yielded an 85.6% sensitivity, a 56.8% specificity, and a 4.2% positive predictive value. The BTMed model calibrated to scan the same number of individuals as the PLCO_{m2012} model demonstrated a sensitivity of 76.0%, specificity of 70.9%, and a 5.5% positive predictive value. This level of sensitivity compares favorably with that of the PLCO_{m2012} model applied to this cohort (70.5%), and specificity was comparable (70.9% for BTMed and 70.8% for PLCO_{m2012}). Dement and colleagues applied the 2013 USPSTF-recommended screening criteria (age 50–80 years, ≥ 30 pack-years of smoking, and quitting < 15 years in past) to their data set and obtained a 50.9% sensitivity, an 81.2% specificity, and a 5.7% positive predictive value.⁸² This large decline in sensitivity, from 85.6% to 50.9%, using the different eligibility criteria, represents a failure of the 2013 USPSTF criteria (which exclude occupation) to detect as many as 40% of the lung cancers detected in the Welch and colleagues study.

5 | RECOMMENDATIONS

5.1 | Pre-eminence of primary prevention of occupational lung cancer

Primary prevention of occupational lung cancer through the control of exposure to lung carcinogens in the workplace is key to avoiding future suffering from occupational lung cancer. Elimination of current and future exposures to occupational lung carcinogens, education of workers, employers and other parties to the workplace, and regulations by governments and authorities play key roles in this form of prevention.

Secondary prevention of occupational lung cancer promotes the avoidance of unnecessary morbidity and mortality through the early detection of the cancer, after occupational exposure to a lung carcinogen has occurred and at a stage at which it is potentially curable. LDCT-based lung cancer screening provides the basis for this

secondary prevention and also provides an excellent opportunity to promote smoking cessation, which is a second form of primary prevention. LDCT screening does not prevent the occurrence of lung cancer but it can prevent lung cancer mortality.

Finally, *tertiary prevention* of occupational lung cancer aims to reduce the consequences of living with lung cancer and its treatment. Recent advancements in the treatment of lung cancer are considerable and translate into longer survival for many people with the disease. Job accommodation is important to allow people with lung cancer to return to work if able and to continue to earn income and thereby lessen the impact of the diagnosis on themselves and their families. Just financial compensation for occupational lung cancer as an occupational disease can also play an invaluable role in softening the impact of a frequently devastating disease for workers and their families.

5.2 | Underlying principles for LDCT in occupational populations

The Collegium Ramazzini recognizes key principles supporting lung cancer screening among workers at risk.

First, the urgency presented by the magnitude of occupational lung cancer in combination with current evidence in favor of lung cancer screening efficacy supports the use of LDCT screening for workers at risk.

Second, in recognition of the principles of respect for autonomy and justice, the Collegium believes that at-risk workers should be provided with a choice about undergoing lung cancer screening. This decision should be made in consultation with their doctors who, together with other members of the medical and scientific community, are duty-bound to provide ample and accessible information to enable good decision-making. Such information should include the limits and risks of annual low dose CT scans, including the possible detection of health conditions for which current treatment is inadequate (e.g., malignant mesothelioma of the pleura). Participation in lung cancer screening should be voluntary.

Third, working populations have often been kept uninformed about their occupational exposures to lung carcinogens. Even if informed, conditions of the workplaces frequently prevented avoidance of such exposures. The identification of resources to support lung cancer screening and subsequent diagnosis and treatment for these workers should be a high priority for unions and other organizations representing them, and for government agencies and employers responsible for a safe and healthy workplace—that is, for primary prevention.

Fourth, additional research to determine the optimal parameters to ensure effective occupational lung cancer screening should be funded and undertaken. Of particular concern is whether non-malignant chronic lung disease associated with asbestos, silica, beryllium, and other lung carcinogens significantly alters the risk-benefit ratio of the use of annual LDCT. In addition, the comorbidities due to smoking may affect screening outcomes in occupational

populations whose eligibility criteria include a lesser tobacco use history in combination with exposure to workplace lung carcinogens. The Collegium encourages research collaboration and data pooling in occupational lung cancer screening studies that include long term mortality follow-up to develop an improved understanding of the effectiveness of screening working populations at increased risk of lung cancer and certain workers within those populations, such as those with pneumoconiosis.

5.3 | Eligibility for LDCT screening

Workers at risk for occupational lung cancer will benefit from determination of eligibility for lung cancer screening that takes into account (a) age and smoking, and (b) a number of other lung cancer risk factors. These include occupational exposure to lung carcinogens, personal history of cancer (excluding skin cancer), personal history of chronic lung disease, family history of lung cancer, and other demonstrated lung cancer risk factors.

The Collegium recommends that persons at risk for lung cancer from occupational exposures be offered annual LDCT if their cumulative risk of lung cancer approximates the level of risk endorsed by the guidelines promulgated by the United States Preventive Services Task Force (USPSTF) in 2021 and the National Comprehensive Cancer Network (NCCN) in the United States in 2021. At present, these agencies recommend screening for people aged 50 and over who have smoked at least 20 pack-years of cigarettes. The Collegium recommends that additional lung cancer risk factors, including exposure to known or suspected occupational and environmental lung carcinogens, family history of lung cancer (especially among first degree relatives and relatives ≤ 60 years of age), a personal history of chronic obstructive lung disease, pneumoconiosis, or pulmonary fibrosis, or a personal history of cancer (excluding skin cancer) be considered as part of the risk assessment for eligibility determination for lung cancer screening. The period of time since initial occupational exposure (latency) is another factor that should be considered (e.g., ≥ 15 years). If the presence of these additional risk factors, in combination with age and smoking history, is associated with a level of risk that meets or exceeds the level of risk identified by the USPSTF and NCCN, then an annual low dose chest CT for lung cancer screening should be offered. We do not favor a specific age cut-off at which to end screening, but we recognize that only persons who are sufficiently healthy and have sufficient life expectancy to undergo diagnostic work-up and potentially curative treatment should be offered screening for lung cancer. In view of the rising risk of occupational lung cancer over time and the potential or actual interaction between occupational lung carcinogens and cigarette smoking even after quitting, screening programs may choose to screen workers with occupational lung cancer risk for prolonged periods after they have quit smoking cigarettes. The Collegium acknowledges that there are uncertainties and assumptions entailed in this approach and that risk assessment for individual workers necessitates application of

significant professional judgment. We encourage the implementation of well-organized screening programs that can further our knowledge about optimal occupation-inclusive lung cancer screening strategies.

It is the responsibility of occupational health and medical professionals and stakeholders (governments, employers, statutory insurance agencies, and labor unions) to identify worker populations that have excess lung cancer risk. Such populations include those with known exposure to known or suspected lung carcinogens or working in occupations/trades or work tasks with known elevated risk for lung cancer. As a general rule, having worked in such conditions for ≥ 5 years constitutes significant risk, though shorter periods of exposure also may be significant, depending on intensity and frequency of exposures.

5.4 | Elements of high-quality occupational lung cancer screening

Lung cancer screening is effective if it is conducted as part of a systematic approach that has the following elements^{62,83}:

1. Eligibility determination, including ability to undergo potentially curative treatment of lung cancer
2. Provision of information on lung cancer risks, risks and benefits of screening, screening process and outcomes, and expected follow-up
3. Informed decision-making
4. Advice on smoking cessation
5. CT scan acquisition with control over quality and radiation dose
6. CT scan interpretation by experienced readers and with use of a standardized reading protocol
7. Follow-up of CT results with appropriate health care providers according to recognized guidelines
8. Access to and plan to involve appropriate medical specialties for work-up and treatment of lung cancer, as well as unanticipated findings that require further medical evaluation.

Ideally, lung cancer screening will be conducted under the auspices of an organized screening program or research study that can provide all of the elements listed above. However, it is unlikely that all workers at risk of occupational lung cancer on a global basis will have access to such programs. And lung cancer screening cannot be effectively restricted to wealthy countries with the resources to establish such dedicated programs. Simplified schemes that can be endorsed by conscientious health care providers that can meet local conditions based on available resources need to be developed. Keys to success will be the physical accessibility of screening sites for at-risk worker populations and implementation of methods that can monitor and assure quality of the screening process.

Advances in lung cancer screening have brought this domain closer to other accepted cancer screening methods such as screening for colon, breast and cervical cancer. As such, in the presence of radiologic expertise in the use of LDCT scans to screen for lung

cancer and pulmonary, oncologic and surgical expertise in the diagnosis and treatment of lung cancer (significant barriers, admittedly), lung cancer screening can become a part of the occupational health service (OHS) or primary health care. Key to success is the OHS or primary health care provider's role in taking a good occupational history to identify work-related risks.

The recommendations contained in this statement are based primarily on a high likelihood of mortality reduction in the screened population and do not address issues of cost-effectiveness.

5.5 | Responsibilities of parties to promote lung cancer screening

Uptake of lung cancer screening in the United States, where low dose CT scanning for this purpose has been approved since 2013, has been disappointingly slow. It currently stands at less than 20% of the people who are eligible for low dose CT scanning based on age and smoking criteria.⁸⁴

The Collegium Ramazzini recognizes that achieving widespread lung cancer screening for workers at risk will be an enormous challenge. The challenges are manifold: organizational, educational, fiscal, regulatory, and political. Even the first step is rarely taken: workers are unaware of their health risks and their health providers evince little or no interest in the health consequences of work.

The Collegium calls upon governments and other stakeholders to undertake organized activities to promote lung cancer screening among workers who are at elevated risk of lung cancer. These activities can include public education campaigns to increase awareness among both workers and health care providers of occupational lung cancer risk and the importance of lung cancer screening; development of lung cancer screening eligibility guidelines and reimbursement policies that address occupational lung cancer risks; initiation or enhancement of mechanisms to identify, educate and motivate occupational populations at increased risk of lung cancer, including use of exposure registries; development of programs with organized labor and employers to encourage use of lung cancer screening; and support of research to understand and apply effective methods of increasing participation in lung cancer screening by blue collar workers.

Developing and implementing a system of workplace-based funded and independently-administered occupational health care—from primary prevention to tertiary prevention—would greatly enhance both the reduction or elimination of exposure to occupational lung carcinogens and the early detection and treatment of lung cancer.

5.6 | Equity challenges within and across countries

The Collegium acknowledges the enormous variation in resources and capacities among and within different countries in provision of health care, including support for cancer screening. Lung cancer

screening involves the use of costly CT scanners and the participation of skilled radiologists, representing a challenge for all countries, especially low- and middle-income countries. These expenses are amplified by the costs and sophisticated medical care associated with the diagnosis and treatment of lung cancer.

This challenge is intensified by the fact that the highest smoking rates in the world are nowadays in low- and middle-income countries, which will therefore face an enormous burden of lung cancer in the future. These countries include especially those in Asia and Eastern Europe.⁸⁵ Many of these same countries likely have widely prevalent exposures to insufficiently controlled occupational lung carcinogens and agents known to be associated with chronic obstructive and fibrotic lung disease.

We note that among the largest lung cancer screening programs in the world has been undertaken by a middle-income country—China.⁸⁶

The Collegium supports efforts to decrease the cost of lung cancer screening through the application of automated artificial intelligence (AI) interpretation of CT scans, development of cost-effective lung cancer screening programs, and the mass utilization of less-expensive CT scanners that can nonetheless obtain images sufficient for lung cancer screening.

More challenging in many parts of the world will be ramping up the human and facility resources of existing health care systems to ensure proper diagnosis and treatment for people identified as having likely lung cancers as part of screening programs. This is no small feat in countries where lung cancer death rates are increasing—low and middle income countries.

In addition, strenuous efforts should be made within countries to ensure an equitable application of lung cancer screening across racial and ethnic groups within occupational populations at risk. The issue is not only a matter of basic equity, but it is likely that many of the workers who have been most highly exposed to occupational lung carcinogens belong to economically deprived racial and ethnic groups.

5.7 | Ethical considerations

The Collegium emphasizes the rights of workers to a safe and healthy working environment free from exposure to any cancer risks. The Collegium Ramazzini takes note of the resolution recently passed by the United Nations Human Rights Council endorsing a “human right to a clean, healthy and sustainable environment.”⁸⁷ In June 2022, the International Labor Organization (ILO) amended the ILO Declaration on Fundamental Principles and Rights at Work to include “a safe and healthy working environment” as a fundamental principle and right at work. However, in the absence of such a clean work environment, workers also have the right to make decisions freely about how they want to mitigate such risks once they have been incurred. These rights are foundational to fair and equitable approaches to occupational health.

The Collegium therefore recognizes the need to advocate for the concurrent introduction of lung cancer screening in current

occupational and primary care practice alongside simultaneous with important systematic efforts to understand, identify and prevent exposures to lung carcinogens in the workplace.

Once introduced, lung cancer screening should only be offered on a voluntary basis with a procedure for ensuring the consent of the participant is fully informed about benefits as well as risks.

Workers should be informed, as in all disease screening efforts, of the risks and benefits of screening for lung cancer. Among such risks (and benefits) are incidental health findings for which there is presently little curative treatment, such as malignant mesothelioma among asbestos-exposed workers.

Further research concerning the effectiveness and the optimal application of lung cancer screening for occupational populations is needed (see Recommendations 2 and 9). However, the need for additional knowledge should not serve as a pretext for inaction in the conduct of lung cancer screening among workers at risk, even as additional research is undertaken.

5.8 | Role of smoking cessation

All lung cancer screening programs should include an evidence-based smoking cessation component. The health benefits of smoking cessation importantly apply to lung cancer risk but additionally have an enormous benefit for the prevention of many types of cancer and other causes of mortality.

5.9 | Research needs

Gaps in knowledge about lung cancer screening in people exposed to workplace risks are considerable, but, as noted above, it should not be cause for delaying its application. It is important to recognize that the question of whether lung cancer screening is beneficial has been answered resoundingly in the affirmative. Lung cancer screening programs, once introduced, provide the basis for addressing many of the items provided they are designed to appropriately collect the needed data.

We acknowledge that limited progress has been made in developing and validating a generalizable lung cancer risk model that include a wide variety of occupational lung cancer risks. We recognize, however, that this important area of cancer prevention is highly dynamic, and the occupational health community is encouraged to both track ongoing research and undertake screening research studies to better understand which workers will benefit from the application of low dose CT scan-based screening. Pooling occupational screening studies with mortality follow-up would be an important step to addressing current important knowledge gaps.

Important research issues remain beyond the critical ones of evaluating LDCT efficacy and effectiveness in screening participants identified via a combination of a smoking history and additional occupational risk factors for lung cancer. They include, at a minimum,

optimizing LDCT and risk factor information to tailor the application of screening; cost-effectiveness; the integration of biomarkers to improve risk stratification⁸⁸ and cancer identification; identifying the best screening intervals; and how best to implement LDCT screening in target occupational populations.⁸⁹ For workers at risk of occupational exposure-related comorbidities, such as asbestosis, silicosis, and chronic obstructive lung disease, it will be important to understand the impact of these comorbidities on the real world effectiveness of lung cancer screening, diagnosis and treatment.

AUTHOR CONTRIBUTIONS

Steven Markowitz, Knut Ringen, and John M. Dement conceived the work and wrote the first draft. All authors edited the Statement and added intellectual contributions. All authors have approved this manuscript version for submission and agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The Collegium Ramazzini conducted peer review of the Statement and its membership approved the final Statement.

ACKNOWLEDGMENTS

This statement has been approved by the membership of the Collegium Ramazzini.

STATEMENT ON OCCUPATIONAL LUNG CANCER SCREENING

The Collegium Ramazzini is an international scientific society that examines critical issues in occupational and environmental medicine with a view towards action to prevent disease and promote health. The Collegium derives its name from Bernardino Ramazzini, the father of occupational medicine, a professor of medicine of the Universities of Modena and Padua in the late 1600s and the early 1700s. The Collegium is comprised of 180 physicians and scientists from 35 countries, each of whom is elected to membership. The Collegium is independent of commercial interests.

CONFLICTS OF INTEREST STATEMENT

Dr. Steven Markowitz has a financial assistance agreement with the U.S. Department of Energy (DOE) to screen DOE workers for lung cancer using low dose chest CT scans. Drs. Knut Ringen and John Dement have a financial assistance agreement with the U.S. Department of Energy (DOE) to screen DOE construction workers for lung cancer using low dose chest CT scans. Dr. Dennis Nowak is a member of an advisory board of Pfizer Inc. regarding reimbursement of varenicline for smoking cessation. Dr. Nowak provides clinical, pharmacological, and psychological support for nicotine abstinence in smokers and high risk workers. The remaining authors declare no conflict of interest.

DISCLOSURE BY AJIM EDITOR OF RECORD

John Meyer declares that he has no conflict of interest in the review and publication decision regarding this article.

DATA AVAILABILITY STATEMENT

Data sharing not applicable to this article as no data sets were generated or analyzed during the current study.

ORCID

Steven Markowitz  <http://orcid.org/0000-0002-0052-6745>

Knut Ringen  <http://orcid.org/0000-0001-5930-3768>

L. Christine Oliver  <http://orcid.org/0000-0002-1866-2163>

Rodney Ehrlich  <http://orcid.org/0000-0001-5736-9237>

REFERENCES

- Compton K, Dean FE, Fu W, et al. Cancer incidence, mortality, years of life lost, years lived with disability, and disability-adjusted life years for 29 cancer groups from 2010 to 2019: a systematic analysis for the global burden of disease study 2019. *JAMA Oncol*. 2022;8(3):420-444. doi:10.1001/jamaoncol.2021.6987
- Bray F, Ferlay J, Soerjomataram I, Siegel RL, Torre LA, Jemal A. Global cancer statistics 2018: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. *CA Cancer J Clin*. 2018;68(6):394-424. doi:10.3322/caac.21492
- Straif K. The burden of occupational cancer. *Occup Environ Med*. 2008;65(12):787-788. doi:10.1136/oem.2007.038224
- Global Burden of Disease 2016 Occupational Carcinogens Collaborators. Global and regional burden of cancer in 2016 arising from occupational exposure to selected carcinogens: a systematic analysis for the Global Burden of Disease Study 2016. *Occup Environ Med*. 2020;77(3):151-159. doi:10.1136/oemed-2019-106012
- Loomis D, Guha N, Hall AL, Straif K. Identifying occupational carcinogens: an update from the IARC monographs. *Occup Environ Med*. 2018;75(8):593-603. doi:10.1136/oemed-2017-1049
- Rushton L, Bagga S, Bevan R, et al. Occupation and cancer in Britain. *Br J Cancer*. 2010;102(9):1428-1437. doi:10.1038/sj.bjc.6605637
- Labrèche F, Kim J, Song C, et al. The current burden of cancer attributable to occupational exposures in Canada. *Prev Med*. 2019;122:128-139. doi:10.1016/j.ypmed.2019.03.016
- Ahn YS, Jeong KS. Epidemiologic characteristics of compensated occupational lung cancers among Korean workers. *J Korean Med Sci*. 2014;29(11):1473-1481. doi:10.3346/jkms.2014.29.11.1473
- Kim EA, Lee HE, Kang SK. Occupational burden of cancer in Korea. *Saf Health Work*. 2010;1(1):61-68. doi:10.5491/SHAW.2010.1.1.61
- Brown T, Darnton A, Fortunato L, Rushton L. Occupational cancer in Britain: respiratory cancer sites: larynx, lung and mesothelioma. *Br J Cancer*. 2012;107(suppl 1):S56-S70. doi:10.1038/bjc.2012.119
- Carder M, Darnton A, Gittins M, et al. Chest physician-reported, work-related, long-latency respiratory disease in Great Britain. *Eur Respir J*. 2017;50(6):1700961. doi:10.1183/13993003.00961-2017
- Del Bianco A, Demers PA. Trends in compensation for deaths from occupational cancer in Canada: a descriptive study. *CMAJ Open*. 2013;1(3):E91-E96. doi:10.9778/cmajo.20130015
- IARC. List of classifications by cancer sites with sufficient or limited evidence in humans, IARC monographs volumes 1-135. 2023. Accessed January 22, 2024. https://monographs.iarc.who.int/wp-content/uploads/2019/07/Classifications_by_cancer_site.pdf. 2023
- Cogliano VJ, Baan R, Straif K, et al. Preventable exposures associated with human cancers. *J Natl Cancer Inst*. 2011;103(24):1827-1839. doi:10.1093/jnci/djr483
- Environmental Protection Agency. *EPA Releases First Major Update to Chemicals List in 40 years*. Environmental Protection Agency; 2019. <https://www.epa.gov/newsreleases/epa-releases-first-major-update-chemicals-list-40-years>
- Kauppinen T. Occupational exposure to carcinogens in the European Union. *Occup Environ Med*. 2000;57(1):10-18. doi:10.1136/oem.57.1.10
- Kauppinen T, Uusitalo S, Saalo A, et al. Trends of occupational exposure to chemical agents in Finland in 1950-2020. *Ann Occup Hyg*. 2013;57(5):593-609. doi:10.1093/annhyg/mes090
- Peters CE, Ge CB, Hall AL, Davies HW, Demers PA. CAREX Canada: an enhanced model for assessing occupational carcinogen exposure. *Occup Environ Med*. 2015;72(1):64-71. doi:10.1136/oemed-2014-102286
- Nicholson WJ, Perkel G, Selikoff IJ. Occupational exposure to asbestos: population at risk and projected mortality—1980-2030. *Am J Ind Med*. 1982;3(3):259-311. doi:10.1002/ajim.4700030305
- Flanagan DM. Asbestos. In: USGS. 2016 Minerals Yearbook May 2018, pp. 10. Accessed January 19, 2024. <https://d9-wret.s3-us-west-2.amazonaws.com/assets/palladium/production/mineral-pubs/asbestos/myb1-2016-asbes.pdf>
- Markowitz SB, Dickens B. Screening for occupational lung cancer. *Clin Chest Med*. 2020;41(4):723-737. doi:10.1016/j.ccm.2020.08.016
- Syamlal G, Jamal A, Mazurek JM. Combustible tobacco and smokeless tobacco use among working adults—United States, 2012 to 2014. *J Occup Environ Med*. 2016;58(12):1185-1189. doi:10.1097/JOM.0000000000000898
- Gonghuan Y, Hsia J, Yan Y, eds. *Global Adult Tobacco Survey (GATS) China 2010 Country Report*. 2010; Chinese Center for Disease Control and Prevention. https://cdn.who.int/media/docs/default-source/ncds/ncd-surveillance/data-reporting/china/gats/en-gats-china-report.pdf?sfvrsn=e58d489d_3&download=true
- Nielsen LS, Bælum J, Rasmussen J, et al. Occupational asbestos exposure and lung cancer—a systematic review of the literature. *Arch Environ Occup Health*. 2014;69(4):191-206. doi:10.1080/19338244.2013.863752
- Markowitz SB, Levin SM, Miller A, Morabia A. Asbestos, asbestosis, smoking, and lung cancer: new findings from the North American insulator cohort. *Am J Respir Crit Care Med*. 2013;188(1):90-96. doi:10.1164/rccm.201302-0257OC
- Wang X, Yano E, Qiu H, et al. A 37-year observation of mortality in Chinese chrysotile asbestos workers. *Thorax*. 2012;67(2):106-110. doi:10.1136/thoraxjnl-2011-200169
- Offermans NSM, Vermeulen R, Burdorf A, et al. Occupational asbestos exposure and risk of pleural mesothelioma, lung cancer, and laryngeal cancer in the prospective Netherlands cohort study. *J Occup Environ Med*. 2014;56(1):6-19. doi:10.1097/JOM.000000000000060
- Olsson AC, Vermeulen R, Schüz J, et al. Exposure-response analyses of asbestos and lung cancer subtypes in a pooled analysis of case-control studies. *Epidemiology*. 2017;28(2):288-299. doi:10.1097/EDE.0000000000000604
- De Matteis S, Consonni D, Lubin JH, et al. Impact of occupational carcinogens on lung cancer risk in a general population. *Int J Epidemiol*. 2012;41(3):711-721. doi:10.1093/ije/dys042
- Liu Y, Steenland K, Rong Y, et al. Exposure-response analysis and risk assessment for lung cancer in relationship to silica exposure: a 44-year cohort study of 34,018 workers. *Am J Epidemiol*. 2013;178(9):1424-1433. doi:10.1093/aje/kwt139
- Consonni D, Matteis SD, Pesatori AC, et al. Lung cancer risk among bricklayers in a pooled analysis of case-control studies. *Int J Cancer*. 2015;136(2):360-371. doi:10.1002/ijc.28986
- Lacourt A, Pintos J, Lavoué J, Richardson L, Siemiatycki J. Lung cancer risk among workers in the construction industry: results from two case-control studies in Montreal. *BMC Public Health*. 2015;15(941):941. doi:10.1186/s12889-015-2237-9
- Kachuri L, Villeneuve PJ, Parent MÉ, Johnson KC, Harris SA. Occupational exposure to crystalline silica and the risk of lung cancer in Canadian men. *Int J Cancer*. 2014;135(1):138-148. doi:10.1002/ijc.28629

34. Silverman DT, Samanic CM, Lubin JH, et al. The diesel exhaust in miners study: a nested case-control study of lung cancer and diesel exhaust. *J Natl Cancer Inst.* 2012;104(11):855-868. doi:10.1093/jnci/djs034
35. Pintos J, Parent ME, Richardson L, Siemiatycki J. Occupational exposure to diesel engine emissions and risk of lung cancer: evidence from two case-control studies in Montreal, Canada. *Occup Environ Med.* 2012;69(11):787-792. doi:10.1136/oemed-2012-100964
36. Schubauer-Berigan MK, Daniels RD, Pinkerton LE. Radon exposure and mortality among white and American Indian uranium miners: an update of the Colorado Plateau cohort. *Am J Epidemiol.* 2009;169(6):718-730. doi:10.1093/aje/kwn406
37. Leuraud K, Schnelzer M, Tomasek L, et al. Radon, smoking and lung cancer risk: results of a joint analysis of three European case-control studies among uranium miners. *Radiat Res.* 2011;176(3):375-387. doi:10.1667/rr2377.1
38. Kreuzer M, Sobotzki C, Schnelzer M, Fenske N. Factors modifying the radon-related lung cancer risk at low exposures and exposure rates among German uranium miners. *Radiat Res.* 2018;189(2):165-176. doi:10.1667/RR14889.1
39. Denholm R, Schüz J, Straif K, et al. Is previous respiratory disease a risk factor for lung cancer. *Am J Respir Crit Care Med.* 2014;190(5):549-559. doi:10.1164/rccm.201402-0338OC
40. Blanc PD, Annesi-Maesano I, Balmes JR, et al. The occupational burden of nonmalignant respiratory diseases. an official American Thoracic Society and European Respiratory Society Statement. *Am J Respir Crit Care Med.* 2019;199(11):1312-1334. doi:10.1164/rccm.201904-0717ST
41. Takiguchi Y. Chronic obstructive pulmonary disease as a risk factor for lung cancer. *World J Clin Oncol.* 2014;5(4):660-666. doi:10.5306/wjco.v5.i4.660
42. Naccache JM, Gibiot Q, Monnet I, et al. Lung cancer and interstitial lung disease: a literature review. *J Thorac Dis.* 2018;10(6):3829-3844. doi:10.21037/jtd.2018.05.75
43. Sone S, Takashima S, Li F, et al. Mass screening for lung cancer with mobile spiral computed tomography scanner. *Lancet.* 1998;351(9111):1242-1245. doi:10.1016/S0140-6736(97)08229-9
44. Henschke CI, McCauley DI, Yankelevitz DF, et al. Early lung cancer action project: overall design and findings from baseline screening. *Lancet.* 1999;354(9173):99-105. doi:10.1016/S0140-6736(99)06093-6
45. Henschke CI, Yankelevitz DF, Libby DM, et al. Survival of patients with stage I lung cancer detected on CT screening. *N Engl J Med.* 2006;355(17):1763-1771. doi:10.1056/NEJMoa060476
46. Aberle DR, Adams AM, National Lung Screening Trial Research Team, et al. Reduced lung-cancer mortality with low-dose computed tomographic screening. *N Engl J Med.* 2011;365(5):395-409. doi:10.1056/NEJMoa1102873
47. de Koning HJ, van der Aalst CM, de Jong PA, et al. Reduced lung-cancer mortality with volume CT screening in a randomized trial. *N Engl J Med.* 2020;382(6):503-513. doi:10.1056/NEJMoa1911793
48. Tammemägi MC, Katki HA, Hocking WG, et al. Selection criteria for lung-cancer screening. *N Engl J Med.* 2013;368(8):728-736. doi:10.1056/NEJMoa1211776
49. Cassidy A, Myles JP, van Tongeren M, et al. The LLP risk model: an individual risk prediction model for lung cancer. *Br J Cancer.* 2008;98(2):270-276. doi:10.1038/sj.bjc.6604158
50. Toumazis I, Bastani M, Han SS, Plevritis SK. Risk-Based lung cancer screening: a systematic review. *Lung Cancer.* 2020;147:154-186. doi:10.1016/j.lungcan.2020.07.007
51. Bach PB, Kattan MW, Thornquist MD, et al. Variations in lung cancer risk among smokers. *J Natl Cancer Inst.* 2003;95(6):470-478. doi:10.1093/jnci/95.6.470
52. ten Haaf K, Jeon J, Tammemägi MC, et al. Risk prediction models for selection of lung cancer screening candidates: a retrospective validation study. *PLoS Med.* 2017;14(4):e1002277. doi:10.1371/journal.pmed.1002277
53. Katki HA, Kovalchik SA, Berg CD, Cheung LC, Chaturvedi AK. Development and validation of risk models to select ever-smokers for CT lung cancer screening. *JAMA.* 2016;315(21):2300-2311. doi:10.1001/jama.2016.6255
54. Tammemägi MC, Ruparel M, Tremblay A, et al. USPSTF2013 versus PLCom2012 lung cancer screening eligibility criteria (International Lung Screening Trial): interim analysis of a prospective cohort study. *Lancet Oncol.* 2022;23(1):138-148. doi:10.1016/S1470-2045(21)00590-8
55. U.S. Preventative Services Task Force. *Final Recommendation Statement, Lung Cancer: Screening.* U.S. Preventative Services Task Force; 2021. <https://www.uspreventiveservicestaskforce.org/uspstf/recommendation/lung-cancer-screening>
56. SAPEA (Science Advice for Policy by European Academies) *Improving Cancer Screening in the European Union.* SAPEA (Science Advice for Policy by European Academies). 2022. <https://sapea.info/topic/cancer-screening>
57. Veronesi G, Baldwin D, Henschke C, et al. Recommendations for implementing lung cancer screening with low-dose computed tomography in Europe. *Cancers.* 2020;12(6):1672. doi:10.3390/cancers12061672
58. Zhou Q, Fan Y, Wang Y, et al. China national lung cancer screening guideline with low-dose computed tomography (2018 version). *Zhongguo fei ai za zhi.* 2018;21(2):67-75. doi:10.3779/j.issn.1009-3419.2018.02.01
59. He J, Li N, Chen WQ, et al. China guideline for the screening and early detection of lung cancer (2021, Beijing). *Zhonghua zhong liu za zhi.* 2021;43(3):243-268. doi:10.3760/cma.j.cn112152-20210119-00060
60. Koegelenberg CFN, Dorfman S, Schewitz I, et al. Recommendations for lung cancer screening in Southern Africa. *J Thorac Dis.* 2019;11(9):3696-3703. doi:10.21037/jtd.2019.08.66
61. Wood DE, Kazerooni EA, Baum SL, et al. Lung cancer screening, version 3.2018, NCCN clinical practice guidelines in oncology. *J Natl Compr Canc Netw.* 2018;16(4):412-441. doi:10.6004/jnccn.2018.0020
62. National Comprehensive Cancer Network Clinical Practice Guidelines in Oncology. *Lung Cancer Screening Version 1.2012 (2012) and Version 1.2022 (October 26, 2021).* 2012/2022; National Comprehensive Cancer Network Clinical Practice Guidelines in Oncology. 2012/2022. https://cdn-links.lww.com/permalink/jti/a/jti_2015_01_09_bartolmai_jti-14-182r1_sdc1.pdf
63. Wolff H, Vehmas T, Oksa P, Rantanen J, Vainio H. Asbestos, asbestosis, and cancer, the Helsinki criteria for diagnosis and attribution 2014: recommendations. *Scand J Work Environ Health.* 2015;41(1):5-15. doi:10.5271/sjweh.3462
64. Delva F, Margery J, Laurent F, Petitprez K, Pairon JC. Medical follow-up of workers exposed to lung carcinogens: French evidence-based and pragmatic recommendations. *BMC Public Health.* 2017;17(1):191-198. doi:10.1186/s12889-017-4114-1
65. Tiitola M, Kivisaari L, Huuskonen MS, et al. Computed tomography screening for lung cancer in asbestos-exposed workers. *Lung Cancer.* 2002;35(1):17-22. doi:10.1016/s0169-5002(01)00294-x
66. Minniti S, Valentini M, Pozzi Mucelli R. Low-dose helical chest CT in asbestos-exposed workers in the veneto region: preliminary results. *Radiol Med (Torino).* 2005;110(4):317-324.
67. Das M, Mühlenbruch G, Mahnken AH, et al. Asbestos surveillance program aachen (ASPA): initial results from baseline screening for lung cancer in asbestos-exposed high-risk individuals using low-dose multidetector-row CT. *Eur Radiol.* 2007;17(5):1193-1199. doi:10.1007/s00330-006-0426-8

68. Fasola G, Belvedere O, Aita M, et al. Low-dose computed tomography screening for lung cancer and pleural mesothelioma in an asbestos-exposed population: baseline results of a prospective, nonrandomized feasibility trial—an Alpe-Adria Thoracic Oncology Multidisciplinary Group study (ATOM 002). *Oncologist*. 2007;12(10):1215-1224. doi:10.1634/theoncologist.12-10-1215
69. Vierikko T, Jarvenpaa R, Autti T, et al. Chest CT screening of asbestos-exposed workers: lung lesions and incidental findings. *Eur Respir J*. 2006;29(1):78-84. doi:10.1183/09031936.00073606
70. Mastrangelo G, Ballarin MN, Bellini E, et al. Feasibility of a screening programme for lung cancer in former asbestos workers. *Occup Med*. 2008;58(3):175-180. doi:10.1093/occmed/kqn018
71. Clin B, Morlais F, Guittet L, et al. Performance of chest radiograph and CT scan for lung cancer screening in asbestos-exposed workers. *Occup Environ Med*. 2009;66(8):529-534. doi:10.1136/oem.2008.041525
72. Kato K, Gemba K, Ashizawa K, et al. Low-dose chest computed tomography screening of subjects exposed to asbestos. *Eur J Radiol*. 2018;101:124-128. doi:10.1016/j.ejrad.2018.02.017
73. Maisonneuve P, Rampinelli C, Bertolotti R, et al. Low-dose computed tomography screening for lung cancer in people with workplace exposure to asbestos. *Lung Cancer*. 2019;131:23-30. doi:10.1016/j.lungcan.2019.03.003
74. Brims FJH, Harris EJA, Murray C, et al. Lung cancer screening an asbestos exposed population: existing lung cancer risk criteria are not sufficient. *Respirology*. 2023;28(6):543-550. doi:10.1111/resp.14487
75. Ollier M, Chamoux A, Naughton G, Pereira B, Dutheil F. Chest CT scan screening for lung cancer in asbestos occupational exposure. *Chest*. 2014;145(6):1339-1346. doi:10.1378/chest.13-2181
76. Roberts HC, Patsios DA, Paul NS, et al. Screening for malignant pleural mesothelioma and lung cancer in individuals with a history of asbestos exposure. *J Thorac Oncol*. 2009;4(5):620-628. doi:10.1097/JTO.0b013e31819f2e0e
77. Barbone F, Barbiero F, Belvedere O, et al. Impact of low-dose computed tomography screening on lung cancer mortality among asbestos-exposed workers. *Int J Epidemiol*. 2018;47(6):1981-1991. doi:10.1093/ije/dyy212
78. Loewen G, Black B, McNew T, Miller A. Lung cancer screening in patients with Libby amphibole disease: high yield despite predominantly environmental and household exposure. *Am J Ind Med*. 2019;62(12):1112-1116. doi:10.1002/ajim.23042
79. Heidrich J, Centmayer A, Wolff C, et al. A nationwide program for lung cancer screening by low-dose CT among formerly asbestos-exposed workers in Germany: concept and participation. *Saf Health Work*. 2022;13:S130-S131.
80. Markowitz SB, Manowitz A, Miller JA, et al. Yield of low-dose computerized tomography screening for lung cancer in high-risk workers: the case of 7,189 US nuclear weapons workers. *Am J Public Health*. 2018;108(10):1296-1302. doi:10.2105/AJPH.2018.304518
81. Welch LS, Dement JM, Cranford K, et al. Early detection of lung cancer in a population at high risk due to occupation and smoking. *Occup Environ Med*. 2019;76:137-142. doi:10.1136/oemed-2018-105431
82. Dement JM, Ringen K, Hines S, Cranford K, Quinn P. Lung cancer mortality among construction workers: implications for early detection. *Occup Environ Med*. 2020;77(4):207-213. doi:10.1136/oemed-2019-106196
83. Wilson JMG, Jungner G. *Principles and Practice of Screening for Disease*. World Health Organization; 1968:168.
84. Yong PC, K, Rehmani S, J, Kale MS, et al. Lung cancer screening uptake in the United States. *Chest*. 2020;157(4):236-238. doi:10.1016/j.chest.2019.08.2176
85. World Population Review 2022. 2022. Accessed January 19, 2024. <https://worldpopulationreview.com/country-rankings/smoking-rates-by-country>
86. Li N, Tan F, Chen W, et al. One-off low-dose CT for lung cancer screening in China: a multicentre, population-based, prospective cohort study. *Lancet Respir Med*. 2022;10(4):378-391. doi:10.1016/S2213-2600(21)00560-9
87. United Nations General Assembly. *Resolution Adopted by the Human Rights Council on 8 October 2021: 48/13 The human right to a clean, healthy, and sustainable environment*. United Nations General Assembly. <https://digitallibrary.un.org/record/3983329>
88. Freitas C, Sousa C, Machado F, et al. The role of liquid biopsy in early diagnosis of lung cancer. *Front Oncol*. 2021;11(11):634316. doi:10.3389/fonc.2021.634316
89. Oudkerk M, Liu S, Heuvelmans MA, Walter JE, Field JK. Lung cancer LDCT screening and mortality reduction—evidence, pitfalls and future perspectives. *Nat Rev Clin Oncol* 2021;18(3):135-151. doi:10.1038/s41571-020-00432-6.

How to cite this article: Markowitz S, Ringen K, Dement JM, et al. Occupational lung cancer screening: a Collegium Ramazzini statement. *Am J Ind Med*. 2024;67:289-303. doi:10.1002/ajim.23572