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Projected Lifetime Cancer Risks From Current Computed Tomography Imaging

Rebecca Smith-Bindman, MD; Philip W. Chu, MS; Hana Azman Firdaus, MPH; Carly Stewart, MHA; Matthew Malekhedayat, BS; Susan Alber, PhD; Wesley E. Bolch, PhD; Malini Mahendra, MD; Amy Berrington de González, DPhil; Diana L. Miglioretti, PhD

IMPORTANCE Approximately 93 million computed tomography (CT) examinations are performed on 62 million patients annually in the United States, and ionizing radiation from CT is a known carcinogen.

OBJECTIVE To project the number of future lifetime cancers in the US population associated with CT imaging in 2023.

DESIGN, SETTING, AND PARTICIPANTS This risk model used a multicenter sample of CT examinations prospectively assembled between January 2018 and December 2020 from the University of California San Francisco International CT Dose Registry. Data analysis was conducted from October 2023 to October 2024.

MAIN OUTCOMES AND MEASURES Distributions of CT examinations and associated organ-specific radiation doses were estimated by patient age, sex, and CT category and scaled to the US population based on the number of examinations in 2023, quantified by the IMV national survey. Lifetime radiation-induced cancer incidence and 90% uncertainty limits (UL) were estimated by age, sex, and CT category using National Cancer Institute software based on the National Research Council's Biological Effects of Ionizing Radiation VII models and projected to the US population using scaled examination counts.

RESULTS An estimated 61 510 000 patients underwent 93 000 000 CT examinations in 2023, including 2 570 000 (4.2%) children, 58 940 000 (95.8%) adults, 32 600 000 (53.0%) female patients, and 28 910 000 (47.0%) male patients. Approximately 103 000 (90% UL, 96 400-109 500) radiation-induced cancers were projected to result from these examinations. Estimated radiation-induced cancer risks were higher in children and adolescents, yet higher CT utilization in adults accounted for most (93 000; 90% UL, 86 900-99 600 [91%]) radiation-induced cancers. The most common cancers were lung cancer (22 400 cases; 90% UL, 20 200-25 000 cases), colon cancer (8700 cases; 90% UL, 7800-9700 cases), leukemia (7900 cases; 90% UL, 6700-9500 cases), and bladder cancer (7100 cases, 90% UL, 6000-8500 cases) overall, while in female patients, breast was second most common (5700 cases; 90% UL, 5000-6500 cases). The largest number of cancers was projected to result from abdomen and pelvis CT in adults, reflecting 37 500 of 103 000 cancers (37%) and 30 million of 93 million CT examinations (32%), followed by chest CT (21500 cancers [21%]; 20 million examinations [21%]). Estimates remained large over a variety of sensitivity analyses, which resulted in a range of 80 000 to 127 000 projected cancers across analyses.

CONCLUSIONS AND RELEVANCE This study found that at current utilization and radiation dose levels, CT examinations in 2023 were projected to result in approximately 103 000 future cancers over the course of the lifetime of exposed patients. If current practices persist, CT-associated cancer could eventually account for 5% of all new cancer diagnoses annually.

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Supplemental content

Author Affiliations: Author affiliations are listed at the end of this article.

Corresponding Author: Rebecca Smith-Bindman, MD, Department of Epidemiology and Biostatistics, University of California, San Francisco, 550 16th St, Box 0560, San Francisco, CA 94143 (rebecca.smith-bindman@ucsf.edu).

omputed tomography (CT) is an indispensable and widely performed medical imaging test. Ongoing technological advancements expand its capabilities and popularity, and utilization continues to rise in the United States, exceeding prepandemic volume. While CT aids diagnosis, leading to improved outcomes, it also exposes patients to ionizing radiation at levels known to be associated with increased cancer risk. Several large retrospective cohort studies have shown that childhood exposure to CT is associated with increased risk of hematologic malignant neoplasms and brain cancer.²⁻⁵ In adults, cancer risks from low to moderate radiation doses are primarily based on studies of Japanese atomic bomb survivors or populations irradiated through medical or occupational exposures.^{6,7} However, there is also evidence that CT damages DNA in adults.8 Radiation-induced cancer risks from CT examinations vary by radiation dose, which depends on the clinical indication; body region imaged; patients' sex, age, and size; and acquisition techniques. 9 A 2009 analysis¹⁰ estimated that approximately 29 000 future cancers would result from routine CT exposures in the United States in 2007. The study authors used best-available data on the volume and distribution of examinations, approximations of radiation doses, and associated absorbed organ doses. Since then, the number of CT examinations performed annually in the United States has increased by more than 30%, 1 more granular data have become available describing examination types, and more accurate methods have been developed for estimating organ dose.

This study updates previously projected lifetime cancer incidence associated with CT using the most recent utilization numbers available, empirical data on CT type by age and sex, and organ doses estimated directly from examination-level clinical data across the United States using best-practice methods. The purpose is to understand the public health impact of current CT use and to identify the highest risk examination types, age, and sex groups.

Methods

This risk model used patient-level data from the University of California San Francisco (UCSF) International CT Dose Registry, which has assembled CT examinations from 143 US hospitals and outpatient facilities associated with 22 health care organizations in 20 states. 9 For each examination, the registry captured Digital Imaging and Communications in Medicine (DICOM) metadata, including patient age, sex, effective diameter of the body part imaged, scanner type, examination name and description, and other technical acquisition parameters, such as kilovoltage, milliamperage, scan length, phase, pitch, and collimation. The UCSF Committee on Human Research approved the study with a waiver of consent due to the large number of records making it impractical to contact all participants, the researchers not knowing the identity of the participants; and the risk of contacting participants being greater than the study risks. Collaborating institutions obtained local ethics approval. We have followed the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) reporting guideline.

Key Points

Question How many future cancers could result from radiation exposure from annual computed tomography (CT) examinations in the United States?

Findings In this risk model, the 93 million CT examinations performed in 62 million patients in 2023 were projected to result in approximately 103 000 future cancers. Although the per-examination cancer risk was higher in children, higher CT utilization among adults accounted for the majority of the projected cancers.

Meaning These findings suggest that if current radiation dosing and utilization practices continue, CT-associated cancers could eventually account for 5% of all new cancer diagnoses annually.

CT Utilization

We used the IMV Medical Information Division CT Market Outlook Report, based on a national, annual survey of 235 hospitals and 78 imaging facilities, to quantify the number of CT examinations performed in the United States in 2023.1 IMV medical imaging utilization data have been validated against sources such as Medicare and the Veterans Administration by the US National Council on Radiation Protection Report No. 184 and used in several publications. 11,12 To apportion examinations between adults and children, we used the proportion of pediatric examinations in 2022 in the American College of Radiology (ACR) National Radiology Data Registry (Judy Burleson, MHSA, and Mike Simanowith, MD, ACR, email, November 13, 2023). To estimate the number of patients who underwent CT in 2023, longitudinal data from the registry from 2016 and 2020 were used to estimate the annual number of examinations per patient by age and sex (mean ranged from 1.1-1.7). This average was applied to the total number of CT examinations in 2023, by sex and age group, to estimate the number of patients exposed.

Distribution of Examinations by Age, Sex, and CT Category

Using DICOM metadata, CT examinations in the registry were assigned to 1 of 26 CT categories that reflect a combination of body region and clinical indication (18 in adults¹³; 13 in children [Denise Bos, MD, unpublished data, March 2025) (eMethods in Supplement 1). Some CT categories represent a single body region (eg, cervical spine), while other regions are subdivided into categories reflecting radiation dose needs of the underlying indication (eg, in the abdomen, low dose includes imaging for kidney stones, routine dose for trauma, and high dose for cancer).

To estimate distributions of scans by age, sex, and CT category, we used pediatric examinations (ages 0-17 years) from the registry from January 2018 to December 2020 (46 559 patients) and adult examinations (ages 18-99 years) from January to December 2020 (74 653 patients). We excluded CT examinations associated with biopsies and procedures, positron emission tomography, or research (all infrequent) as well as age, sex, and CT category strata with fewer than 12 examinations given that estimated doses could be imprecise. Additional years

of data were used for pediatric examinations (2018-2020) compared with adult examinations (2020) to ensure stable estimates within strata. We verified that the distribution of CT categories and radiation dose per category remained stable in adults from 2018 to 2020. From this sample of 121 212 examinations, we estimated the proportions of examinations by age, sex, and CT category resulting in 418 strata: 288 in adults (18 CT categories, 8 age groups, and 2 sexes); and 130 in children (13 CT categories, 5 age groups, 2 sexes).

Individual Patient-Dependent Organ Dose Reconstruction

We estimated absorbed doses (radiation transport code MCNPX version 2.70 [Los Alamos National Laboratory]) for 18 organs for each CT examination through Monte Carlo radiation transport simulations using exact, examination-level technical parameters and patient size mapped to morphometry-matched hybrid computational phantoms from the University of Florida/National Cancer Institute phantom library. We then calculated mean organ doses (and SDs) in milliGray (mGy) for each strata.

Statistical Analysis

Cancer Risk Estimation

We projected future lifetime radiation-induced cancer risk using the National Cancer Institute's Radiation Risk Assessment Tool (RadRAT)^{18,19} software version 4.3.1, which utilizes risk models from the National Academy of Sciences' Biologic Effects of Ionizing Radiation (BEIR) VII report for 11 site-specific cancers (stomach, colon, liver, lung, breast, uterus, ovary, prostate, bladder, and thyroid cancer and leukemia), plus 7 additional cancer sites (oral cavity or pharynx, esophagus, rectum, pancreas, kidney, and brain or central nervous system cancer plus a remainder category) using a more recent follow-up of the Japanese atomic bomb survivors and pooled analyses of other medically exposed cohorts. 18 For a given cancer type, RadRAT estimates excess lifetime risk of cancer from the time of exposure based on user-supplied organ dose and US life table estimates of age- and sex-specific baseline cancer rates. These risk estimates account for death as a competing risk using sex-specific life table estimates for the US 2019 population. We developed solutions to expedite bulk use of RadRAT to estimate risks within the 418 strata (eMethods in Supplement 1).

Cancer Projections

We scaled the registry-based distribution of CT categories by age and sex by the IMV-derived total number of examinations, using the ACR percentage of pediatric examinations, to estimate the distribution of examinations at the US population level in 2023. We excluded examinations that occurred in the last year of life, which are unlikely to contribute to future cancers given the average latency between CT exposure and radiation-induced cancer development. To determine this proportion, we quantified the number of CT examinations performed in 2022 in patients' last 1 and 2 years of life for each strata of age and sex at Kaiser Permanente Northern California, following published methods. ^{19,20} Overall, 10.6% of scans were performed in the last year of life (9.4% in female pa-

tients and 12.1% in male patients), varying from 0.9% in children ages 1 to 4 years (1.4% girls and 0.5% boys) to 38.6% in adults ages 90 to 99 years (35.1% women and 44.4% men). We then applied the projected cancer rates from RadRAT to nationally scaled examination counts (reduced by the proportion of end-of-life examinations) to estimate lifetime cancer incidence and 90% uncertainty limits (UL) resulting from CT examinations in 2023. Since future cancer estimates are based on a linear model of the total radiation dose received, the projected number of cancers remains the same regardless of whether the analysis is based on the number of patients (62 million, who each underwent an average of 1.5 scans) or examinations (93 million).

Uncertainty Estimates and Sensitivity Considerations

RadRAT uses Monte Carlo simulation based on Latin hypercube sampling to account for uncertainty in the radiation risk model coefficients, transfer of risks from the Japanese to the US population, the dose and dose rate reduction effectiveness factor (DDREF), uncertainty in organ doses, and adjustments to minimal latency periods. 18 A latency adjustment was phased in between 4.0 and 11.0 years after exposure for solid cancers, 0.4 and 4.1 years for leukemia, and 2.5 and 7.6 years for thyroid cancer. To represent uncertainty in the adjustments for minimum latency on risk estimates, the midpoint, μ, is described by the following triangular probability distributions: solid cancers other than thyroid, T(5, 7.5, 10); thyroid, T(3, 5, 7); and leukemia, T(2, 2.25, 2.5), where numbers represent time after exposure in years. RadRAT outputs 90% ULs, providing an upper and lower estimate of potential future cancers.

Sensitivity analyses were conducted modifying the baseline model assumptions. First, we applied male lung cancer risk coefficients to female patients in our projections because some epidemiological studies have not supported the 3-fold higher risk of radiation-induced lung cancer in female compared with male patients in BEIR VII.²¹ Second, we reduced the estimated annual imaging volume by 10% to account for potential overestimation by IMV, and third, we increased it by 10% for potential underestimation. Fourth, we reduced organ doses by 20% to allow for possible national differences from the UCSF registry, and fifth, we increased them by 20%. Sixth, we applied the higher IMV estimate of the percentage of CT examinations in children (9.0% vs 3.3%). Seventh, we used the distribution of examinations by age and CT category from calendar years 2018 to 2019, in case the 2020 distribution was atypical due to COVID-19. Last, we excluded CT examinations performed in the last 2 years, rather than 1 year, of life, varying by age and sex. All analyses used SAS version 9.3 (SAS Institute) and R version 4.2.2 (2022-10-31 ucrt [R Project for Statistical Computing]). Data analysis was conducted from October 2023 to October 2024.

Results

Ninety-three million CT examinations were performed in 61510000 patients in the United States in 2023, including an

estimated 3 069 000 CTs (3.3%) in 2 570 000 children (4.2%) and 89 931 000 CTs (96.7%) in 58 940 000 adults (95.8%) (Table 1; eTable 1 in Supplement 1). Patients underwent a mean of 1.5 examinations each, varying by age (Table 1), and the median number of examinations per patient was 1 across all age groups. The total number of examinations increased with age for all CT categories, peaking in adults ages 60 to 69 years (Figure 1; eTable 1 in Supplement 1). After excluding examinations performed in the last year of life, a total of 84 161 000 were included for estimating cancer risks.

Organ Doses

Organ doses by body regions and sex are shown for sample age strata (eTable 2 in Supplement 1). Doses were similar but not identical by sex for most categories. For example, the mean (SD) brain dose for routine-dose head CT in children ages 5 to 9 years was 5% higher in boys (48.0 [27.3] mGy) than in girls (45.7 [24.1] mGy). Other categories, such as full body, had larger differences. For example, there was a 29% increase in pancreas dose between boys aged 5 to 9 years (21.5 [13.5] mGy) vs girls aged 5 to 9 years (16.7 [8.9] mGy). In general, organ doses were similar in children and adults or increased with age. For example, the mean (SD) colon dose in routine abdomen and pelvis CT was approximately twice as high in women aged 50 to 59 years (25.4 [15.2] mGy) vs girls aged 5 to 9 years (12.8 [8.7] mGy). However, there were exceptions: organ doses were highest overall in children younger than 1 year (eg, mean [SD] brain dose for routine head CT in boys <1 year was 60.0 [36.5] mGy), and mean (SD) bone marrow doses in head CT decreased with age (eg, boys <1 year, 26.7 [16.7] mGy; boys aged 5-9 years, 14.6 [10.0] mGy; and men aged 50-59 years, 3.5 [2.7] mGy).

Projected Cancer Risks

CT utilization in the United States in 2023 was estimated to result in 102700 (90% UL, 96 400-109 500) projected lifetime cancers, including 93 000 (90% UL, 86 900-99 600) in adults and 9700 (90% UL, 8100-11 600) in children (Table 2). The leading cancers in adults were lung cancer (21 400 [90% UL, 19 200-24 000]), colon cancer (8400 [90% UL, 7500-9400]), and leukemia (7400 [90% UL, 6100-8900]), whereas the most frequent projected cancers in children were thyroid (3500 [90% UL, 2300-5500]), lung (990 [90% UL, 870-1100]), and breast (630 [90% UL, 550-730]) cancer (Table 2; eFigure in Supplement 1). Lung and thyroid cancer incidence were higher in female patients, whereas incidence of most other cancers was similar by sex or slightly higher in male patients (eFigure and eTable 3 in Supplement 1).

Projected Cancers by Age

Projected cancer risks per CT examination were estimated to be highest among children who underwent CT at younger than 1 year and decreased with age at exposure (Figure 1). For example, cancer risk in girls younger than 1 year were 20 cancers per 1000 examinations (1900 of 97 000) versus 2 per 1000 in girls aged 15 to 17 years (1100 of 483 600) (eTables 1 and 4 in Supplement 1). However, despite the higher risk per examination in children, higher utilization contributed to more projected cancers in adults (Table 2 and Figure 2). CT use in

adults aged 50 to 59 years was associated with the highest number of projected cancers: 10 400 (90% UL, 8200-13 000) in female patients and 9300 (90% UL, 7500-11700) in male patients (eTable 4 in Supplement 1).

Projected Cancers by CT Category

Abdomen and pelvis CT was estimated to contribute the largest number of projected cancers (40%) in adults (37 500 [90% UL, 32 900-42 600] cases), whereas head CT contributed the largest number of cancers (53%) in children (5100 [90% UL, 3700-7100) (Table 2 and **Figure 3**; eTable 3 in Supplement 1). For a few categories, such as full body, the projected proportion of cancers (8000 [7.8%]) was greater than the proportion of scans (4 607 000 scans [5.0%]) (Tables 1 and 2).

Sensitivity Analyses

The sensitivity analyses generated a range in estimated future cancers from 79 900 to 126 600 by reducing and increasing organ doses by 20%, respectively, reflecting 22.2% fewer cancers to as many as 23.3% more cancers than the primary analysis (eTable 6 in Supplement 1). Using the IMV-estimated proportion of pediatric examinations resulted in an 11.0% increase in projected cancers overall (to 114 000) and an increase in the proportion of cancers from childhood imaging from 9.4% to 23.2%.

Discussion

CT is frequently lifesaving, yet its potential harms are often overlooked, and even very small cancer risks will lead to a significant number of future cancers given the tremendous volume of CT use in the United States. For current utilization and radiation dosing practices, we projected approximately 103 000 future cancers could result from CT use in the United States in 2023 (with sensitivity analyses projecting a range of 80 000 to 127 000) among the 62 million people who underwent CT. To provide context, if the number of new cancer diagnoses in the United States remains stable (1.95 million in 2023) and both the utilization and radiation doses from CT remain unchanged in future decades, CT could be responsible for approximately 5% of cancers diagnosed each year. This would place CT on par with other significant risk factors, such as alcohol consumption (5.4%) and excess body weight (7.6%).

The projected number of radiation-induced cancers in this analysis is 3 to 4 times higher than the earlier assessment of CT exposure for several reasons. ¹⁰ First, while growth in utilization has slowed over the intervening years, ²⁰ CT use is 30% higher today than in 2007, due to growth in low-value, potentially unnecessary imaging ²³⁻²⁷ as well as population aging. Second, dose modeling in this study accounted for multiphase scanning, which occurs in 28.5% of examinations but was not modeled in the prior study, as multiphase frequency was unknown. Third, the substantially higher organ doses in this study were reconstructed using newer dosimetry methods with examination-level data from more than 120 000 actual examinations, while the prior study modeled doses from national survey data or imaging protocols and assumed examinations in

Table 1. Estim	ated Number of P	Table 1. Estimated Number of Patients Who Underwent CT and Number of CT Examinations in the United States in 2023, by Sex, Age, and Body Region ^a	nt CT and Number o	of CT Examinatio	ns in the United	States in 2023, b	y Sex, Age, and E	ody Region ^a			
			Fxaminations	CT examination type, No.	type, No.						
Age, y	Patients, No.	CT examinations, No. (%)	per patient, mean, No. ^b	Abdomen and pelvis	Head	Chest	Spine	Head and neck combined	Cardiac	Full body ^c	Extremities ^d
Total population											
All	61 510 000	93 000 000 (100)	1.51	30 221 400	24115800	19975300	7 735 200	2 8 2 6 9 0 0	1073900	4 607 000	2 444 500
Child	2570000	3 069 000 (3.3)	1.19	006689	1 600 800	275 200	290 000	NA	26 400	57 300	129 300
Adult	58 940 000	89931000 (96.7)	1.53	29 531 500	22 515 000	19700100	7 445 200	2826900	1047500	4 5 4 9 7 0 0	2315200
Female											
All ages	32 600 000	48 549 200 (52.2)	1.49	16 669 900	12 263 700	10136700	4 082 800	1494100	513 800	2 2 2 8 0 0 0	1160200
~	000 06	97 000 (0.1)	1.12	4000	72 200	13100	3100	NA	4500	0	0
1-9	370 000	439300 (0.5)	1.18	83 100	255800	43 300	38 500	NA	3200	7900	7400
10-17	740 000	870 000 (0.9)	1.18	270600	370500	73 100	93 500	NA	4000	14 500	43 900
18-39	2 9 6 0 0 0 0 0	7 972 500 (8.6)	1.34	3 676 800	1 948 000	1117300	651300	190 200	0	219 100	169 800
40-59	8 980 000	13147100(14.1)	1.46	5 136 000	2 953 300	2 590 900	1 004 100	328 700	244 400	289 900	299 800
62-09	12 020 000	18767100(20.2)	1.56	5 546 600	4 290 900	5 01 0 8 0 0	1 600 000	504 500	237 200	1059500	517 700
80-99	4 4 4 0 0 0 0 0	7 256 200 (7.8)	1.63	1 952 800	2 373 000	1288200	692 300	470 700	20 500	337 100	121 600
Male											
All ages	28 910 000	44 450 900 (47.8)	1.54	13 551 400	11852300	9838700	3 652 500	1332800	560 100	2379100	1284500
<1	100 000	119800(0.1)	1.14	6300	87 000	17 400	2200	NA	2900	1100	0
1-9	490 000	580 100 (0.6)	1.18	108 700	356400	45 200	48 900	NA	3500	10 800	0029
10-17	780 000	962 800 (1.0)	1.23	217 100	459 000	83 100	103 800	NA	5300	23 100	7400
18-39	5 160 000	7 282 700 (7.8)	1.41	2 501 800	2 070 800	000 606	754900	278 100	29 000	344 300	364 800
40-59	7 7 7 0 0 0 0 0	11780600(12.7)	1.52	3 928 500	2 872 600	2 458 500	931900	304 600	249 200	645 300	390 100
62-09	11 320 000	18278300 (19.7)	1.61	5 414 100	4 327 000	2 080 600	1 336 400	456 300	237 200	1059500	367 200

^d Extremities includes all upper and lower extremity examinations. More granular results by CT category within each body region are in eTable 1 in Supplement 1. Row values may

^c Full body includes whole body examinations in children and combined chest, abdomen, and pelvis examinations

293 800

1244900

1 679 500

84300

295 000

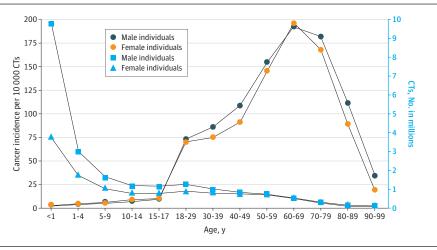
not sum to "CT examinations" column due to rounding.

Abbreviations: CT, computed tomography; NA, not applicable, meaning this category does not exist in this

1.66 1.61

62-09 80-99 age group.

Figure 1. Number of Computed Tomography (CT) Examinations and Cancer Incidence by Sex



The projected number of future cancers (left axis; dark blue and orange circles) was estimated using the reduced number of CT examinations (excluding examinations that occur in the last year of life) as reported in Table 2. Cancer incidence was based on the total number of examinations (right axis; light blue circles and triangles), a conservative estimate.

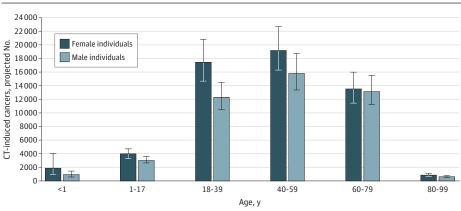
Table 2. Projected Number of Future Cancers Overall and by Cancer Type Associated With CT Examinations Performed in the United States in 2023, by Age Group and Body Region^a

	Projected future cancers, No. (90% UL)			
Cancer type	All CT examinations (N = 93 000 000)	CT examinations in adults (n = 89 931 000)	CT examinations in children (n = 3 069 000)	
Total	102 700 (96 400-109 500)	93 000 (86 900-99 600)	9700 (8100-11600)	
Projected cancer by type				
Lung	22 400 (20 200-25 000)	21 400 (19 200-24 000)	990 (870-1100)	
Colon	8700 (7800-9700)	8400 (7500-9400)	330 (280-390)	
Leukemia	7900 (6700-9500)	7400 (6100-8900)	550 (380-820)	
Bladder	7100 (6000-8500)	6900 (5700-8200)	250 (200-320)	
Stomach	7100 (5500-9100)	6800 (5200-8800)	280 (200-400)	
Thyroid	7000 (5400-9200)	3500 (2700-4600)	3500 (2300-5500)	
Breast	5700 (5000-6500)	5100 (4400-5900)	630 (550-730)	
Liver	4100 (3400-5000)	4000 (3200-4900)	160 (130-200)	
Kidney	3000 (2300-3900)	2900 (2200-3700)	130 (90-180)	
Pancreas	2800 (2300-3500)	2700 (2200-3400)	100 (80-140)	
Oral cavity or pharynx	2800 (2300-3400)	2300 (1900-2900)	450 (310-650)	
Brain or CNS	1600 (1300-2000)	1200 (910-1500)	440 (320-620)	
Esophagus	1500 (1300-1800)	1400 (1200-1700)	110 (90-150)	
Prostate	1500 (820-2700)	1400 (760-2700)	70 (30-170)	
Ovary	890 (670-1200)	850 (630-1100)	40 (30-70)	
Rectum	560 (480-660)	540 (450-630)	30 (20-40)	
Uterus	550 (400-760)	530 (380-730)	30 (16-50)	
Other and ill-defined sites	17 400 (15 300-19 800)	15 800 (13 700-18 200)	1600 (1200-2000)	
Projected cancer by CT examination body region				
Abdomen and pelvis	39 100 (34 600-44 200)	37 500 (32 900-42 600)	1600 (1300-2000)	
Chest	22 700 (19 600-26 300)	21 500 (18 400-25 200)	1200 (960-1400)	
Spine	12 900 (11 500-14 500)	11 600 (10 200-13 200)	1300 (1000-1600)	
Head	12 500 (10 600-14 700)	7300 (6200-8700)	5100 (3700-7100)	
Full body	8000 (7000-9100)	7600 (6600-8800)	320 (260-390)	
Head and neck combined	4100 (3500-4800)	4100 (3500-4800)	NA	
Cardiac	3400 (3200-3700)	3300 (3000-3600)	170 (140-210)	
Extremity	80 (60-90)	70 (50-80)	9 (7-11)	

Abbreviations: CNS, central nervous system; CT, computed tomography; NA, not applicable, meaning this category does not exist in this age group; UL, uncertainty limit.

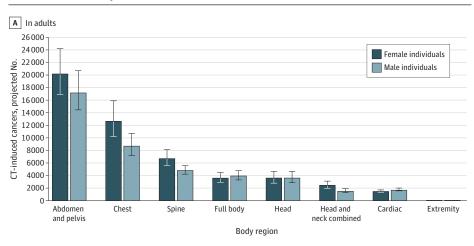
^a More granular results by sex and cross-tabulation by body region and cancer type appear in eTable 3 in Supplement 1.

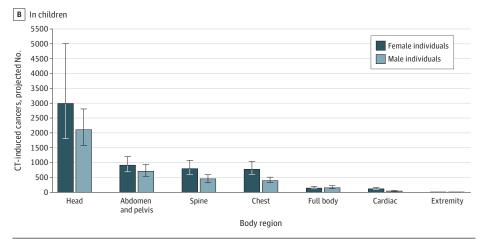
Figure 2. Total Projected Lifetime Cancers by Sex and Age at Exposure



Error bars represent 90% uncertainty limits. CT indicates computed tomography.

Figure 3. Projected Number of Computed Tomography (CT)-Induced Cancers by Body Region Imaged in Adults and Children, by Sex





Error bars represent 90% uncertainty limits.

children were performed using pediatric-specific settings. Lastly, we included more granular CT categories reflecting imaging indications that have important dose differences. Both studies used the same BEIR VII risk models; thus, this would not explain the large observed differences.

Lung cancer was projected to be the most common radiation-induced cancer, with 22 400 cases (eTable 4 in Supple-

ment 1). Approximately 70% of these were in female patients, reflecting the higher BEIR VII risk coefficients in female patients. However, even when we applied male risk coefficients for female examinations, lung cancers were still the most common radiation-induced cancer in female patients. Colon cancer was the next most common, with 8700 cases (58.6% in males). It is unclear whether the current, unexplained in-

crease in these 2 cancers as well as others at unexpectedly younger ages²⁸ may be partly due to CT. Thyroid cancer also revealed notable sex differences, likely due to risk coefficients. For example, 1400 vs 320 thyroid cancers were projected to result from CT exposure in female and male patients, respectively, performed when the patient was younger than 1 year, despite equal thyroid organ doses (74.4 and 75.2 mGy) (eTables 2 and 4 in Supplement 1) and more examinations in male patients (Table 1). Our estimates from childhood CT exposure are higher than those in the large EPI-CT cohort study of pediatric cancer outcomes^{2,3} because we estimated lifetime risk of all cancer types, while EPI-CT examined brain and hematologic cancers within 15 years after exposure.

Abdomen and pelvis CT were projected to cause the greatest number of cancers. These and other examination types, such as high-dose abdomen and pelvis, full body, and spine CT, incur greater risks on average per examination because they frequently use multiple scan phases that result in higher doses. ^{29,30} Often these examinations could use single-phase scanning, which would lower doses without impacting diagnostic accuracy.

This study estimated future cancers using the National Academy of Science BEIR VII-based modeling approach, which is widely accepted in the field of radiation epidemiology. While observational studies have directly quantified childhood cancer risk related to pediatric CT, 2-5 for adult exposures, direct estimates are currently unavailable. To empirically quantify lifetime risk would require decades-long follow-up studies of very large populations, as the Life Span Study has done in Japanese bombing survivors. Thus, to feasibly capture full lifetime risk requires a modeling approach, and there is increasing evidence of elevated cancer risks from other low-dose radiation research supporting these risk estimates. 6,7,31 While the BEIR VII cancer risk models are the most widely used and accepted approach for quantifying the cancer risks from lowdose radiation, several other studies have published risk models, such as the US Environmental Protection Agency, the UK National Radiological Protection Board, and the United Nations Scientific Committee on the Health Effects of Atomic Radiation (UNSCEAR). The risk estimates from these studies are broadly consistent with BEIR VII as well as estimates from the CT study cohorts including EPI-CT.³²

Many of the model assumptions were conservative. For example, we used the ACR's percentage of estimated examinations in children, which is lower than the percentage from IMV. We did not include CT-guided procedures, such as biopsies, which often use higher doses. Furthermore, it is possible that the low-energy x-rays emitted by CT scans cause more cellular damage compared with gamma rays, which were the primary source of radiation released from the atomic bombs.³³ Lastly, RadRAT applied the DDREF of 1.5 (90% uncertainty in-

terval, 1.1-2.3) recommended by BEIR VII to account for differences between low-dose exposure and the higher doses for which the models were developed. This assumes lower radiation doses are less harmful (per unit) than higher doses, based on the BEIR VII estimate that the risk of solid cancer per unit of radiation dose may be 1.5 times lower for doses of 100 mGy or less. However, several systematic reviews of low dose (<100 mGy) and low dose rate exposures support a DDREF of 1. A4,35 If accurate, the true number of projected cancers would be closer to the upper end of the sensitivity estimates than the primary analysis projects.

Strengths and Limitations

This study has several strengths, including detailed data on CT utilization and associated radiation dose, detailed calculation of risks with uncertainty limits, and sensitivity analyses that provide a range of estimates under widely varying assumptions. There are several limitations: first, the BEIR VII risk estimated model parameters are based primarily on the Japanese survivor outcomes, and questions remain about the transfer of radiation risks from the mid-20th century Japanese population to the current US population. The use of a weighted average of the excess relative and excess absolute risk models aims to partly account for this, but these weights are subjective.³⁶ Second, our risk calculations factored in average life expectancies, and the degree to which patients who undergo CT have shorter life expectancy due to underlying illness may overestimate future cancer risk. However, we excluded on average 10.6% of CTs that were likely performed during the last year of life, given these patients are not at risk of a radiation-induced cancer. A recent analysis found that 9.6% of patients who undergo CT died within 1 year,³⁷ similar to our estimate. Third, while the CT categorization algorithm was 90% accurate compared with expert review,¹³ some examinations in the registry may have been miscategorized; however, this is unlikely to significantly impact our results.

Conclusions

In this study, approximately 5% of annual cancer diagnoses or 100 000 cancers were projected to result from CT utilization in 2023. Despite public attention to the potential adverse effects, CT use has grown significantly in the United States since 2009. In 2023, 93 million CT examinations were performed in the United States; in 2007, the number was 68.7 million—a 35% increase incompletely explained by population growth. ³⁸ Justification of use and optimization of dose, including consideration of the need for multiphase examinations, are the tenets of CT imaging and must be applied uncompromisingly to mitigate potential harm.

ARTICLE INFORMATION

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Author Affiliations: Department of Epidemiology and Biostatistics, University of California, San Francisco (Smith-Bindman, Chu, Azman Firdaus, Stewart, Malekhedayat); Department of Obstetrics, Gynecology and Reproductive Sciences, University of California, San Francisco (Smith-Bindman); Philip R. Lee Institute for Health Policy Studies, University of California, San Francisco (Smith-Bindman, Mahendra); Department of Public Health Sciences, University of California, Davis (Alber, Miglioretti); J. Crayton Pruitt Family Department of Biomedical Engineering, University of Florida, Gainesville (Bolch); Division of Pediatric Critical Care, Department of Pediatrics, UCSF Benioff Children's Hospital, University of California, San Francisco (Mahendra); Division of Genetics and Epidemiology, The Institute of Cancer Research, London, United Kingdom (Berrington de González); Kaiser Permanente Washington Health Research Institute, Kaiser Permanente Washington, Seattle (Miglioretti).

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Acquisition, analysis, or interpretation of data: All authors.

Drafting of the manuscript: Smith-Bindman, Chu, Stewart.

Critical review of the manuscript for important intellectual content: All authors.
Statistical analysis: Chu, Alber, Mahendra, Berrington de González, Miglioretti.
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Smith-Bindman, Azman Firdaus, Stewart,
Malekhedayat, Bolch.
Supervision: Smith-Bindman, Chu, Azman Firdaus,
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Dr Smith-Bindman reported being a cofounder of Alara Imaging Inc, a company focused on improving the clinical and operational aspects of health systems, including collecting and reporting radiation dose and image quality associated with computed tomography as part of payer-led quality programs. Alara Imaging played no role in any aspect of the article, and this work does not overlap with Alara's commercial activities. Dr Miglioretti reported receiving grants from the National Institutes of Health during the conduct of the study. No other disclosures were reported.

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REFERENCES

- 1. IMV. 2023 CT market outlook report. Accessed March 10, 2025. https://imvinfo.com/product/2023-ct-market-outlook-report/
- 2. Bosch de Basea Gomez M, Thierry-Chef I, Harbron R, et al. Risk of hematological malignancies from CT radiation exposure in children, adolescents and young adults. *Nat Med.* 2023;29(12):3111-3119. doi:10.1038/s41591-023-02620-0
- 3. Hauptmann M, Byrnes G, Cardis E, et al. Brain cancer after radiation exposure from CT examinations of children and young adults: results from the EPI-CT cohort study. *Lancet Oncol.* 2023;24(1):45-53. doi:10.1016/S1470-2045 (22)00655-6
- 4. Pearce MS, Salotti JA, Little MP, et al. Radiation exposure from CT scans in childhood and subsequent risk of leukaemia and brain tumours: a retrospective cohort study. *Lancet*. 2012;380(9840):499-505. doi:10.1016/S0140-6736 (12)60815-0
- 5. Mathews JD, Forsythe AV, Brady Z, et al. Cancer risk in 680,000 people exposed to computed tomography scans in childhood or adolescence: data linkage study of 11 million Australians. *BMJ*. 2013;346:f2360. doi:10.1136/bmj.f2360
- **6.** National Research Council; Committee to Assess Health Risks from Exposure to Low Level of Ionizing Radiation. *Health Risks From Exposure to Low Levels of Ionizing Radiation: BEIR VII Phase* **2.** National Academies Press: 2006.
- 7. Hauptmann M, Daniels RD, Cardis E, et al. Epidemiological studies of low-dose ionizing radiation and Cancer: Summary Bias Assessment and Meta-Analysis. *J Natl Cancer Inst Monogr.* 2020;2020(56):188-200. doi:10.1093/jncimonographs/lgaa010
- **8**. Nguyen PK, Lee WH, Li YF, et al. Assessment of the radiation effects of cardiac CT angiography using protein and genetic biomarkers. *JACC Cardiovasc Imaging*. 2015;8(8):873-884. doi:10.1016/j.jcmg.2015.04.016
- **9**. Smith-Bindman R, Wang Y, Chu P, et al. International variation in radiation dose for computed tomography examinations: prospective cohort study. *BMJ*. 2019;364:k4931. doi:10.1136/bmj.k4931
- **10**. Berrington de González A, Mahesh M, Kim KP, et al. Projected cancer risks from computed tomographic scans performed in the United States in 2007. *Arch Intern Med*. 2009;169(22):2071-2077. doi:10.1001/archinternmed.2009.440
- 11. Mettler FA Jr, Mahesh M, Bhargavan-Chatfield M, et al. Patient exposure from radiologic and nuclear medicine procedures in the United States: procedure

- volume and effective dose for the period 2006-2016. *Radiology*. 2020;295(2):418-427. doi:10.1148/radiol.2020192256
- 12. Mahesh M, Ansari AJ, Mettler FA Jr. Patient exposure from radiologic and nuclear medicine procedures in the United States and worldwide: 2009-2018. *Radiology*. 2023;307(1):e221263. doi:10.1148/radiol.221263
- **13.** Smith-Bindman R, Yu S, Wang Y, et al. An image quality-informed framework for CT characterization. *Radiology*. 2022;302(2):380-389. doi:10.1148/radiol.2021210591
- **14**. Lee C, Lodwick D, Hurtado J, Pafundi D, Williams JL, Bolch WE. The UF family of reference hybrid phantoms for computational radiation dosimetry. *Phys Med Biol*. 2010;55(2):339-363. doi:10.1088/0031-9155/55/2/002
- **15.** Geyer AM, O'Reilly S, Lee C, Long DJ, Bolch WE. The UF/NCI family of hybrid computational phantoms representing the current US population of male and female children, adolescents, and adults-application to CT dosimetry. *Phys Med Biol.* 2014;59(18):5225-5242. doi:10.1088/0031-9155/59/18/5225
- **16**. Stepusin EJ, Long DJ, Ficarrotta KR, Hintenlang DE, Bolch WE. Physical validation of a Monte Carlo-based, phantom-derived approach to computed tomography organ dosimetry under tube current modulation. *Med Phys.* 2017;44(10): 5423-5432. doi:10.1002/mp.12461
- 17. Long DJ, Lee C, Tien C, et al. Monte Carlo simulations of adult and pediatric computed tomography exams: validation studies of organ doses with physical phantoms. *Med Phys.* 2013;40 (1):013901. doi:10.1118/1.4771934
- 18. National Cancer Institute Division of Cancer Epidemiology and Genetics. Radiation risk assessment tool—lifetime cancer risk from ionizing radiation. Updated February 2024. Accessed January 10, 2025. https://radiationcalculators.cancer.gov/radrat/
- 19. Kwan ML, Miglioretti DL, Bowles EJA, et al. Quantifying cancer risk from exposures to medical imaging in the Risk of Pediatric and Adolescent Cancer Associated with Medical Imaging (RIC) Study: research methods and cohort profile. Cancer Causes Control. 2022;33(5):711-726. doi:10.1007/s10552-022-01556-z
- **20**. Smith-Bindman R, Kwan ML, Marlow EC, et al. Trends in use of medical imaging in US health care systems and in Ontario, Canada, 2000-2016. *JAMA*. 2019;322(9):843-856. doi:10.1001/jama.2019.11456
- 21. Cahoon EK, Preston DL, Pierce DA, et al. Lung, laryngeal and other respiratory cancer incidence among Japanese atomic bomb survivors: an updated analysis from 1958 through 2009. *Radiat Res.* 2017;187(5):538-548. doi:10.1667/RR14583.1
- **22.** Islami F, Marlow EC, Thomson B, et al. Proportion and number of cancer cases and deaths attributable to potentially modifiable risk factors in the United States, 2019. *CA Cancer J Clin*. 2024; 74(5):405-432. doi:10.3322/caac.21858
- 23. Hendee WR, Becker GJ, Borgstede JP, et al. Addressing overutilization in medical imaging. *Radiology*. 2010;257(1):240-245. doi:10.1148/radiol.
- **24**. Schroeder AR, Duncan JR. Overuse of medical imaging and its radiation exposure: who's minding

- our children? *JAMA Pediatr*. 2016;170(11):1037-1038. doi:10.1001/jamapediatrics.2016.2147
- **25**. Chow S, McWilliams A, Kaplan DM, Stephens JR. Things we do for no reason: neuroimaging for hospitalized patients with delirium. *J Hosp Med*. 2019;14(7):441-444. doi:10.12788/jhm.3167
- **26.** Tung M, Sharma R, Hinson JS, Nothelle S, Pannikottu J, Segal JB. Factors associated with imaging overuse in the emergency department: a systematic review. *Am J Emerg Med.* 2018;36(2): 301-309. doi:10.1016/j.ajem.2017.10.049
- 27. Smith M, Saunders R, Stuckhardt L, McGinnis JM, eds; The Institute of Medicine of The National Academies. Best Care at Lower Cost: The Path to Continuously Learning Health Care in America. National Academies of Sciences, Engineering, and Medicine; 2013. Accessed April 6, 2019. http://iom.nationalacademies.org/Reports/2012/Best-Care-at-Lower-Cost-The-Path-to-Continuously-Learning-Health-Care-in-America.
- **28**. Siegel RL, Giaquinto AN, Jemal A. Cancer statistics, 2024. *CA Cancer J Clin*. 2024;74(1):12-49. doi:10.3322/caac.21820
- **29**. Guite KM, Hinshaw JL, Ranallo FN, Lindstrom MJ, Lee FT Jr. Ionizing radiation in

- abdominal CT: unindicated multiphase scans are an important source of medically unnecessary exposure. *J Am Coll Radiol*. 2011;8(11):756-761. doi:10.1016/j.jacr.2011.05.011
- **30**. Johnson PT, Bello JA, Chatfield MB, et al. New ACR Choosing Wisely recommendations: judicious use of multiphase abdominal CT protocols. *J Am Coll Radiol*. 2019;16(1):56-60. doi:10.1016/j.jacr.2018.07.026
- **31.** Berrington de Gonzalez A, Daniels RD, Cardis E, et al. Epidemiological studies of low-dose ionizing radiation and cancer: rationale and framework for the monograph and overview of eligible studies. *J Natl Cancer Inst Monogr*. 2020;2020(56):97-113. doi:10.1093/jncimonographs/lgaa009
- **32**. United Nations Scientific Committee on the Health Effects of Atomic Radiation (UNSCEAR). Sources, effects, and risks of ionizing radiation: report to the General Assembly with scientific annexes. January 2017. Accessed March 10, 2025. https://www.unscear.org/docs/publications/2016/UNSCEAR_2016_GA-Report.pdf
- **33**. Russ E, Davis CM, Slaven JE, Bradfield DT, Selwyn RG, Day RM. Comparison of the medical uses and cellular effects of high and low linear energy transfer radiation. *Toxics*. 2022;10(10):628. doi:10.3390/toxics10100628

- **34.** Shore R, Walsh L, Azizova T, Rühm W. Risk of solid cancer in low dose-rate radiation epidemiological studies and the dose-rate effectiveness factor. *Int J Radiat Biol.* 2017;93(10): 1064-1078. doi:10.1080/09553002.2017.1319090
- **35.** Little MP, Pawel DJ, Abalo K, Hauptmann M. Methodological improvements to meta-analysis of low dose rate studies and derivation of dose and dose-rate effectiveness factors. *Radiat Environ Biophys*. 2021;60(3):485-491. doi:10.1007/s00411-021-00921-x
- **36**. Berrington de Gonzalez A, Iulian Apostoaei A, Veiga LH, et al. RadRAT: a radiation risk assessment tool for lifetime cancer risk projection. *J Radiol Prot*. 2012;32(3):205-222. doi:10.1088/0952-4746/32/3/205
- **37**. Mataac MT, Li X, Rehani MM. What proportion of CT scan patients are alive or deceased after 10 years? *Eur J Radiol*. 2024;178:111629. doi:10.1016/j.ejrad.2024.111629
- **38**. National Council on Radiation Protection and Measurements. Report No. 184—Medical Radiation Exposure of Patients in the United States. Accessed March 10, 2025. https://ncrponline.org/shop/reports/report-no-184-medical-radiation-exposure-of-patients-in-the-united-states-2019/