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### ► To cite this version:

Irina Guseva Canu, Alan Gaillen-Guedy, Ahti Antilla, Sandrine Charles, Sandrine Fraize-Frontier, et al.. Lung cancer mortality in the European cohort of titanium dioxide workers: a reanalysis of the exposure–response relationship. *Occupational and Environmental Medicine*, 2022, 79 (9), pp.637-640. 10.1136/oemed-2021-108030 . anses-03663925

**HAL Id: anses-03663925**

**<https://anses.hal.science/anses-03663925>**

Submitted on 28 Mar 2024

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1 **Lung cancer mortality in the European cohort of titanium dioxide workers: a re-analysis of**  
2 **the exposure-response relationship**

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5 Pukkala<sup>2,8</sup>, Mary Schubauer-Berigan<sup>9</sup>, Kurt Straif<sup>10,11</sup>, Pascal Wild<sup>1</sup>, David B. Richardson<sup>12</sup>

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5 Running Head: Titanium dioxide exposure and lung cancer

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14 on Cancer / World Health Organization, the authors alone are responsible for the views expressed  
15 in this article and they do not necessarily represent the decisions, policy or views of the  
16 International Agency for Research on Cancer / World Health Organization.

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1 ABSTRACT

2 Objectives: Animal bioassays have demonstrated convincing evidence of the potential  
3 carcinogenicity to humans of titanium dioxide (TiO<sub>2</sub>), but limitations in cohort studies have been  
4 identified, among which is the healthy worker survivor effect (HWSE). We aimed to address this  
5 bias in a pooled study of four cohorts of TiO<sub>2</sub> workers.

6 Methods: We re-analysed data on respirable TiO<sub>2</sub> dust exposure and lung cancer mortality among  
7 7341 male workers employed in TiO<sub>2</sub> production in Finland, France, United Kingdom and Italy  
8 using the parametric g-formula, considering three hypothetical interventions: setting annual  
9 exposures at 2.4 (U.S. occupational exposure limit), 0.3 (German limit), and 0 mg/m<sup>3</sup> for 25 and  
10 35 years.

11 Results: The HWSE was evidenced. Taking this into account, we observed a positive association  
12 between lagged cumulative exposure to TiO<sub>2</sub> and lung cancer mortality. The estimated number of  
13 lung cancer deaths at each age group decreased across increasingly stringent intervention levels.  
14 At age 70 years, the estimated number of lung cancer deaths expected in the cohort after 35-year  
15 exposure was 293 for exposure set at 2.4 mg/m<sup>3</sup>, 235 for exposure set at 0.3 mg/m<sup>3</sup>, and 211 for  
16 exposure set at 0 mg/m<sup>3</sup>.

17 Conclusion: This analysis shows that HWSE can hide an exposure-response relationship. It also  
18 shows that TiO<sub>2</sub> epidemiological data could demonstrate an exposure-effects relationship if  
19 analysed appropriately. More epidemiological studies and similar re-analyses of existing cohort  
20 studies are warranted to corroborate the human carcinogenicity of TiO<sub>2</sub>. This human evidence,  
21 when combined with the animal evidence, strengthens the overall evidence of carcinogenicity of  
22 TiO<sub>2</sub>.

23 KEY WORDS

1 Occupational exposure; g-formula; causal inference; retrospective cohort

## 2 INTRODUCTION

3 There is substantial interest in human evidence regarding the carcinogenicity of titanium dioxide  
4 ( $\text{TiO}_2$ ), an odorless white pigment and opacifying agent widely used since the 1920s.  $\text{TiO}_2$  is  
5 classified as possibly carcinogenic to humans by the International Agency for Research on Cancer  
6 based on sufficient evidence from cancer bioassay studies and inadequate evidence from human  
7 cancer studies. Since 2020,  $\text{TiO}_2$  is also classified as a suspected human carcinogen by inhalation  
8 in the European Union according to Regulation n°1272/2008.

9 Prior large cohorts of  $\text{TiO}_2$  workers reported increased mortality from lung cancer, but failed to  
10 observe an exposure–response relationship with cumulative exposure to  $\text{TiO}_2$ ,[1] except in a  
11 French cohort of  $\text{TiO}_2$  workers.[2] One of the key limitations noted in the occupational cohorts  
12 was the potential healthy worker survivor effect (HWSE) which can mask association between  
13 cumulative exposure and lung cancer mortality.[3] For instance, in the pooled European study of  
14  $\text{TiO}_2$  workers, a key study on this topic, no evidence of an association between respirable  $\text{TiO}_2$   
15 exposure and lung cancer mortality was observed despite the excess of lung cancer mortality  
16 among male  $\text{TiO}_2$  workers as compared to the general population (standardised mortality  
17 ratio=1.23; 95% confidence interval (95% CI)=1.10-1.38).[4]

18 In this study, we re-analysed a subset of the pooled European cohort of  $\text{TiO}_2$  workers,[4] to  
19 examine the evidence of HWSE and the exposure-response relationship between cumulative  
20 exposure to  $\text{TiO}_2$  and lung cancer mortality. We implemented the g-computation algorithm  
21 formula (g-formula) recommended for statistical analysis of cohort data in the presence of time-  
22 varying confounders affected by prior exposure, typical of HWSE.[5] The g-formula has been  
23 proven to be an essential method for estimating human health effects of exposures and

1 interventions on exposures in such scenarios.[5] Therefore, we applied it to assess the effect of  
2 three hypothetical interventions of TiO<sub>2</sub> exposure limitation.

### 3 METHODS

#### 4 *Study population*

5 The original pooled European cohort included workers who had been employed at least 1 month  
6 in one of 11 TiO<sub>2</sub> production factories in six European countries (Finland, France, Italy, Norway,  
7 Germany, and the UK).[4] All female workers and male workers with missing death certificates  
8 and/or lacking quantitative exposure estimates were excluded from the exposure-response analysis  
9 of this cohort.[4] We used the same criteria of worker inclusion/exclusion as in original cohort,  
10 but restricted the study to four countries (Finland, France, Italy, and the UK), for which data were  
11 still available and ethical approvals obtained.

#### 12 *Exposure assessment*

13 The factories produced mainly pigment-grade TiO<sub>2</sub>, although TiO<sub>2</sub> form (e.g., particle size and  
14 crystalline phase) was ignored.[6] Information on demographic and employment characteristics  
15 was collected from factories' records describing date of birth, sex, race, and dates of hire, job or  
16 department change, and termination. Estimated cumulative occupational exposure to respirable  
17 TiO<sub>2</sub> dust was derived from job title and work history.[6] Exposure assessments were carried out  
18 at the level of occupational titles for each plant for discrete time periods throughout the history of  
19 plant operations. Lists of occupational titles were compiled and coded for each factory. Exposure  
20 measurement data were obtained from company files along with information such as the area(s) of  
21 the plant where measurements were made, the presence of any local ventilation, the type of the  
22 materials being handled and the purpose of sampling. These were then linked to the work history  
23 of each individual in the cohort to provide exposure estimates.

1 *Statistical analysis*

2 The lengths of follow-up varied between countries and ranged from 1950–1972 until 1997–2001.

3 The primary outcome of interest was death for which the underlying cause was attributed to  
4 cancers of the trachea, bronchus and lung (ICD-9 code 162).

5 A data tabulation of person-periods and events was constructed with one record for each person-  
6 year of observation from date of entry into the analysis until end of follow-up or administrative  
7 censoring of workers alive at age 90 years. Using the observed data, we fitted logistic regression  
8 models for the probability of the outcome of interest, for the probability of remaining at work, and  
9 for the probability of dying from a competing cause, as a function of covariates and estimated  
10 exposure. The cumulative TiO<sub>2</sub> exposure was 10-year lagged (Supplementary material Figure S1  
11 and Technical Appendix 1).

12 The g-formula was implemented by a Monte Carlo simulation based on the regression model  
13 estimates of the probability of termination of employment and death.[5, 7] Ten Monte Carlo  
14 samples per exposure scenario were drawn randomly from the observed cohort and the estimated  
15 parameters from the parametric models to recreate the study data for each person in the sample  
16 under specified exposure intervention. Three hypothetical interventions were considered: setting  
17 workers' annual exposures to 2.4 mg/m<sup>3</sup>, 0.3 mg/m<sup>3</sup> (the currently recommended TiO<sub>2</sub>  
18 occupational exposure limits in the US [8] and Germany [9], respectively), and zero exposure. For  
19 each intervention, we assumed two possible exposure durations: 25 and 35 years and estimated the  
20 expected lung cancer mortality at 60, 70, 80, and 90 years of age. The associated 95%-CIs were  
21 calculated using bootstrap samples.

22 RESULTS

1 The cohort included 7341 workers (Table S2). At the end of the follow-up, 139 lung cancer deaths  
2 were observed. The presence of the HWSE was evident (Figure S1). Being in employment reduced  
3 the risk of lung cancer mortality (OR=0.14, 95%CI=0.08-0.22) and the probability of leaving the  
4 employment increased as a function of TiO<sub>2</sub> exposure (OR=3.55 95%CI=2.82-4.46 ). The OR of  
5 lung cancer death associated with lagged cumulative exposure to TiO<sub>2</sub> was estimated at 1.03 per  
6 1 mg/m<sup>3</sup>-year (95%CI=0.99-1.07), after adjustment for the employment status in previous and  
7 current years and employment duration.

8 G-estimates of lung cancer mortality, derived under the three hypothetical interventions, are shown  
9 in Table 1. The estimated number of lung cancer deaths at 60, 70, 80, and 90 years of age all  
10 decrease across the three interventions considered and for both exposure durations.

## 11 DISCUSSION

12 This re-analysis provides the first evidence of an exposure-response relationship between TiO<sub>2</sub>  
13 cumulative exposure and lung cancer mortality using the parametric g-formula. Adjustment of a  
14 standard regression model for employment status orexposure duration is not sufficient for  
15 complete HWSE correction.[3] However, rank ordering of lung cancer deaths across levels of the  
16 intervention estimated by g-formula is consistent with a positive exposure-response association  
17 between TiO<sub>2</sub> and lung cancer (3% per 1 mg/m<sup>3</sup>-year of respiratory TiO<sub>2</sub>).

18 A limitation of the g-formula is the g-null paradox. The g-formula may be guaranteed some degree  
19 of model misspecification if there is treatment-confounder feedback and the sharp causal null  
20 hypothesis (i.e., the intervention has no effect on any individual's outcome at any time) holds.[10]

21 To address concern that model misspecification may lead to bias, we followed recent guidance  
22 [11] to avoid overly parsimonious models for the components of the g-formula in this analysis.



1 A causal interpretation of the findings requires certain statistical assumptions, including  
2 consistency, positivity, exchangeability.[12] The consistency assumption may be challenging  
3 given the complexity of historical exposure conditions and the various TiO<sub>2</sub> forms, the latter being  
4 insufficiently documented. We nevertheless believe that exposure contrasts are defined here well  
5 enough to support meaningful inference regarding TiO<sub>2</sub>'s effect. The positivity hypothesis (i.e.  
6 observations on exposed and unexposed workers through covariate levels) was difficult to confirm  
7 as 82% of workers were exposed. However, the positive exposure-response relationship based on  
8 continuous cumulative lagged exposure to TiO<sub>2</sub> supports these assumptions.

9 The exchangeability assumption (i.e., no unmeasured confounding) is challenging given the  
10 limitations of available data on smoking. The most complete smoking data were available for the  
11 French cohort and showed no effect of adjustment for smoking on estimates of TiO<sub>2</sub>-lung cancer  
12 mortality associations.[2] Moreover, we were able to assess some other occupational co-exposures  
13 that are suspected or known lung carcinogens, including asbestos, welding fumes, and other  
14 mineral dusts; adjustment for these exposures had no effect on the association with TiO<sub>2</sub>.

15 A confounder typically of concern in occupational cohort mortality studies is the HWSE.[3] In the  
16 current study, this source of confounding was addressed by the g-formula. In prior analyses of  
17 these data using standard multivariable regression method, the association was likely masked by  
18 the HWSE and potentially the exposure misclassification in the early years of follow-up. Prior to  
19 undertaking the g-formula analysis, we assessed the most important HWSE components in these  
20 data and confirmed their presence. Moreover, prior reports of SMR analyses indicated patterns  
21 consistent with the HWSE[3] and in the analysis of the French cohort, duration of employment  
22 was negatively associated with lung cancer mortality.[2] These underscore the relevance of g-  
23 methods in these cohorts and future investigations of TiO<sub>2</sub>'s effects on human health.

1 The estimates with overlapping CI are likely due to the limited statistical power in this study,  
2 which might be addressed by additional follow-up of these cohorts. More epidemiological studies  
3 and similar re-analyses of updated existing cohort studies are warranted to corroborate the human  
4 carcinogenicity of TiO<sub>2</sub>. This human evidence, when combined with the animal evidence,  
5 strengthens the overall evidence of carcinogenicity of TiO<sub>2</sub>.

#### 6 ACKNOWLEDGEMENTS

7 None.

#### 8 CONTRIBUTORS

9 IGC designed and conducted this study and drafted the manuscript, AG-G, DBR, and PW  
10 conducted statistical analyses. KS and MSB centralised the data, facilitated data access, and  
11 obtained IARC ethical approval. SC, SFF, and CM contributed to study coordination and French  
12 ethical approval. All authors discussed the study methodology, read the manuscript, critically  
13 reviewed it and agreed on the final version.

#### 14 FUNDING

15 This work was supported by the ANSES/IST Number 2017-CRD-18.

#### 16 COMPETING INTERESTS

17 None declared.

#### 18 PATIENT CONSENT FOR PUBLICATION

19 Not required.

#### 20 ETHICS APPROVAL

21 The study was authorised by the IARC ethics Committee (IEC project number 18-32). The use of  
22 the data for the French cohort was approved by the French Data Protection Authority (CNIL),  
23 Authorisation No 999250. The use of the data for the UK cohort was approved by the EPS ethics

1 committee of Heriot-Watt University (approval number 19/EA/JC/1). The use of the data for the  
2 Italian cohort was approved by the Comitato Etico Interaziendale A.S.L Citta di Torino (approval  
3 number CS2/1250). The use of the data for the Finnish cohort was approved by Statistics Finland  
4 (approval number TK-53-988-19).

5

## 6 DATA AVAILABILITY STATEMENT

7 Data may be obtained from a third party and are not publicly available.

8

## 9 KEY MESSAGES

### 10 **What is already known about this subject?**

11 Titanium dioxide (TiO<sub>2</sub>) is classified by the International Agency for Research on Cancer as  
12 possibly carcinogenic to humans. In 2020, the European Chemicals Agency classified TiO<sub>2</sub> under  
13 all forms as suspected human carcinogen by inhalation. Prior large cohorts of TiO<sub>2</sub> workers  
14 reported increased mortality from lung cancer but failed to observe an exposure–response  
15 relationship with cumulative exposure to TiO<sub>2</sub>, except in a French cohort of TiO<sub>2</sub> workers. A  
16 concern of potential healthy worker survivor effect (HWSE) has been raised.

### 17 **What are the new findings?**

18 We re-analysed data from the European cohort of TiO<sub>2</sub> workers and found strong evidence of the  
19 HWSE. Taking this into account, a positive exposure–response relationship with 10-years lagged  
20 cumulative exposure to TiO<sub>2</sub> was observed. The estimated number of lung cancer deaths at each  
21 age group decreased across increasingly stringent exposure limits.

### 22 **How might this impact on policy or clinical practice in the foreseeable future?**

23 This analysis shows that HWSE can hide exposure–response relationship. It also shows that TiO<sub>2</sub>  
24 epidemiological data could demonstrate an exposure–effects relationship if analysed appropriately.

1 This human evidence, when combined with the animal evidence, strengthens the overall evidence  
2 of carcinogenicity of TiO<sub>2</sub>.

3

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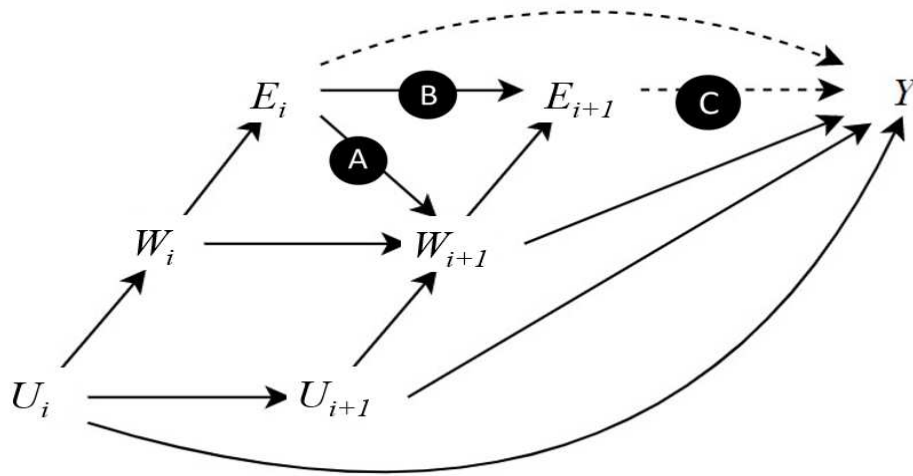
1 Table 1. Estimated cumulative lung cancer mortality at age 60, 70, 80 and 90 years under three  
 2 hypothetical interventions on the TiO<sub>2</sub> exposure and assuming two durations of this exposure.  
 3 European TiO<sub>2</sub> worker cohort, 1955-1990.

4

Attained age (in years)	Cumulative number of lung cancer deaths (95%CI) assuming 25-year exposure			Cumulative number of lung cancer deaths (95%CI) assuming 35-year exposure		
	Set		Set	Set		Set
	exposure to 2.4 mg/m <sup>3</sup>	exposure to 0.3 mg/m <sup>3</sup>	exposure to 0.0 mg/m <sup>3</sup>	exposure to 2.4 mg/m <sup>3</sup>	exposure to 0.3 mg/m <sup>3</sup>	exposure to 0.0 mg/m <sup>3</sup>
60	50 (46-55)	38 (33-44)	39 (34-45)	98 (94-102)	79 (72-86)	74 (68-80)
70	134 (130-139)	105 (97-112)	95 (86-104)	293 (288-299)	235 (226-243)	211 (200-223)
80	168 (162-174)	128 (121-136)	116 (107-125)	417 (407-427)	329 (320-337)	299 (281-316)
90	169 (163-174)	129 (121-136)	116 (108-125)	423 (412-433)	333 (325-342)	303 (286-321)

5  
6

1 ONLINE SUPPLEMENTARY MATERIAL



2

3 Figure S1. Directed acyclic graph representing structural relationships underlying the healthy  
4 worker survivor effect.

5

6 This graph represents the evolution over a worker's lifetime of his or her employment status (1 if  
7 employed, 0 if terminated), exposure and mortality from lung cancer (1 if the cause of death is  
8 lung cancer, 0 for any other situation) as well as the presumed causal links between these factors.

9 Dashed lines show causal effect of the exposure. A, B, and C represent relationships necessary to

10 model for implementation of the g-formula to control for potential confounding bias due to the

11 healthy worker survival effect.  $E_i$  corresponds to exposure to  $\text{TiO}_2$  in year  $i$ ;  $W_i$  corresponds to

12 employment status and  $U_i$  to health status in year  $i$ . Finally,  $Y$  denotes mortality from lung cancer.

13 The unmeasured health status  $U_i$  in year  $i$  influences employment status in the same year and health

14 status  $U_{i+1}$  in the next year. Similarly,  $W_i$  influences  $E_i$  and  $W_{i+1}$ . Finally,  $E_i$  affects  $E_{i+1}$ .

15 Note that this is a simplified directed acyclic graph showing only two time intervals and omitting

16 the baseline covariates.

1 **Technical Appendix 1.**

2

3 Given the objectives of the project, the choice of the g-formula seemed the most judicious, as it  
4 is the only method allowing the evaluation of the effect of hypothetical interventions. To apply  
5 this method we began by modeling the different presumed causal associations presented in Figure  
6 S1: the association between employment status and survival, the association between previous  
7 exposure to TiO<sub>2</sub> and current employment status. The third key component association (i.e., an  
8 association between employment status and subsequent exposure) is considered deterministic,  
9 because the end of employment leads to the end of exposure. For this, the following models were  
10 applied to the observed data in the study population (pooled cohort of four countries, n=7341):

- 11 1. A logistic regression model to predict the probability of job termination for a given person-  
12 year ( $W_{i+1}$  as a function of  $W_i$  and  $E_i$ );
- 13 2. A logistic regression model to predict exposure in that year ( $E_{i+1}$  as a function of  $W_i$  and  $E_i$   
14 when  $W_{i+1} = 1$ ) and a linear regression model to predict exposure level (as a function of  
15 active employment and exposure);
- 16 3. A logistic regression model to predict lung cancer death as a function of employment status  
17 and exposure ( $Y$  as a function of  $E_i$ );
- 18 4. A logistic regression model for other causes of death.

19 The beta coefficients of all variables included in each of these models are provided in the  
20 technical report of the study<sup>1</sup>, available upon request to the corresponding author.

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<sup>1</sup> Irina Guseva Canu, Alan Gaillen-Guedy, Ahti Anttila, Danièle Luce, Damien McElvenny, Franco Merletti, Eero Pukkala, Mary Schubauer-Berigan, Kurt Straif, Pascal Wild, David B. Richardson. Rapport des résultats du projet scientifique « RealyTi » Reanalysis of human TiO<sub>2</sub> data. Troisième version. 22 décembre 2020. 66p.

1 In a second step, we used the results of these models to estimate the potential effect of different  
2 exposure scenarios. The results of this step were expressed in terms of estimated number of deaths  
3 per attained age for each hypothetical intervention on the exposure.

4 We considered three hypothetical interventions:

5 - A first intervention sets the annual respiratory  $\text{TiO}_2$  exposure at  $2.4 \text{ mg/m}^3$

6 - A second intervention sets the annual respiratory  $\text{TiO}_2$  exposure at  $0.3 \text{ mg/m}^3$

7 - A third intervention sets the exposures of all subjects during their entire career to zero

8 These first two values correspond to the currently recommended OELs for  $\text{TiO}_2$  fine dust  
9 (respirable fraction) in the USA and Germany, respectively.

10 The occupational careers, dates of death and dates of lung cancer death of the cohort participants  
11 were resampled (Monte Carlo) 10 times for each scenario, resulting in 10 pseudo-cohorts for each  
12 scenario.

13 In practice, we retained information on the baseline variables for each participant (identification  
14 number, year of hire, age at hire, exposure at the start of follow-up, center identification code in  
15 each country) and drew, year by year, exposure status, cumulative exposure level fixed by  
16 exposure scenario lung cancer death (yes/no), and, if applicable, death by another cause of death  
17 (concurrent causes).

18 Specifically, from year  $m=1$ , employment status is assigned using the conditional probability,  
19 estimated from the parametric employment status model. If the person is employed, the exposure  
20 of this person is defined at the specified level by a new simulation. Then, the probability of lung  
21 cancer and competing causes of death is estimated based on the joint distribution of exposure and  
22 covariates. A binary indicator for each outcome is drawn from a Bernoulli distribution with the



1 associated probability. If the individual is still alive at the end of year  $m$ , the process is repeated  
2 for  $m + 1$ , until death or the end of follow-up date. Within each person-year, we assume the  
3 temporal order of the component variables assigned to each person-year as follows: fixed  
4 covariates, employment status, conditional  $\text{TiO}_2$  exposure, lung cancer death, and death from  
5 another cause.

6 The final step was to calculate for each simulated joint pseudo-cohort the number of deaths in  
7 each age group assuming 25 and 35-year exposure.

8 In order to select the most relevant explanatory variables for each of the four models explained  
9 above, we based ourselves on *Akaike Information Criterion* (AIC). This criterion is based on a  
10 compromise between the quality of the fit and the complexity of the model. This penalizes models  
11 with a large number of variables to limit the effects of over-fitting. The model with the lowest  
12 AIC value was therefore considered the best for predicting employment status, exposure status,  
13 cumulative exposure, and lung cancer death and death from other causes in year  $i$  respectively.

14  
15 The final risk model for lung cancer death for each subject in each year is as follows:

$$\begin{aligned} 17 \quad & \text{logit}(\text{cancer}) \\ 18 \quad & = \beta_1 \text{age} + \beta_2 \text{age}^2 + \beta_3 \text{age}^2 \mathbb{1}_{\text{age} > 65} + \beta_4 \text{age}^2 \mathbb{1}_{\text{age} > 70} \\ 19 \quad & + \beta_5 (\text{year of hire} - 1960)^2 + \beta_6 \text{cumulative exposure}_{\text{lag } 10} \\ 20 \quad & + \beta_7 (\text{cumulative exposure}_{\text{lag } 10})^2 + \beta_8 \mathbb{1}_{\text{active employment}} \\ 21 \quad & + \beta_9 \mathbb{1}_{\text{active employment}_{\text{lag } 1}} + \beta_{10} \text{employment duration} \\ 22 \quad & + \text{intercept} \end{aligned}$$

1 with the  $\beta$  coefficients of each variable calculated for the entire cohort of four countries.  
2 Table S1 confirms that the selected model simulates well the natural course in the joint cohort.  
3 The cumulative exposure variable Lag10 is the lagged cumulative exposure of 10 years. The  
4 application of lag on the cumulative exposure variable consists of introducing a lag time between  
5 exposure and cancer occurrence when modeling the relationship between the two. In practice, this  
6 involves deducting from the cumulative exposure variable the annual exposures over the last 10  
7 years preceding the occurrence of cancer. The 10-year value is used by convention for solid  
8 cancers, although it is possible to derive other, more specific values for a given cancer by  
9 modelling.

10  
11

1 Table S1. Observed and simulated cumulative deaths due to lung cancer by attained age

2

Age (years)	Number of observed death	Number of deaths (associated 95-% confidence interval) estimated by baseline model	
50	9	10	(8-12)
60	43	44	(40-49)
70	99	113	(105-120)
80	138	139	(133-146)
90	139	140	(133-146)

3

4

1 Table S2. Characteristics of the cohort of male titanium dioxide workers

2

<b>Characteristics</b>	<i>N (%)</i>
Number of workers	7341 (100)
Number of workers exposed to TiO <sub>2</sub>	6019 (82.0)
Vital status at the end of follow-up	
Alive	5945 (81.0)
Deceased	1223 (16.7)
From lung cancer	139 (11.4)
From unknown cause	13 (1.1)
Lost to follow-up	173 (2.4)
Number of person-years	179040 (100)
<b>Characteristics</b>	<i>Mean (Standard deviation)</i>
Attained age at hire (years)	28.5 (9.2)
Attained age at start of follow-up (years)	30.2 (9.3)
Attained age at end of follow-up (years)	52.1 (10.9)
Employment duration (years)	13.6 (10.1)
Exposure to TiO <sub>2</sub> (respiratory fraction)*	<i>Mean (Standard deviation) [Min-Max]</i>
Cumulative (mg/m <sup>3</sup> -years)	6.17 (10.47) [0.0003-142.8]
Annual average (mg/m <sup>3</sup> )	0.47 (0.79) [0.00001-8.0]

3 \*Exposure concentration among exposed workers

4

5

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