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Inquinamento atmosferico e mutamenti climatici:
ruolo del medico imparziale ma non neutrale

20, 21 e 22 Ottobre 2023

Auditorium Sede Direzionale Aboca - Loc. Aboca,20 - 52037 Sansepolcro (AR)

Effetti respiratori dell'inquinamento atmosferico e aggiornamento del documento GARD-Italia del 2019

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2005-06 President European Respiratory Society (ERS)

2017-22 WHO - GARD Planning Group Member



Durata: 15'



92-99% of the world population overexposed with respect to the 2021 WHO Air Quality Guidelines



EurRespir J 2021



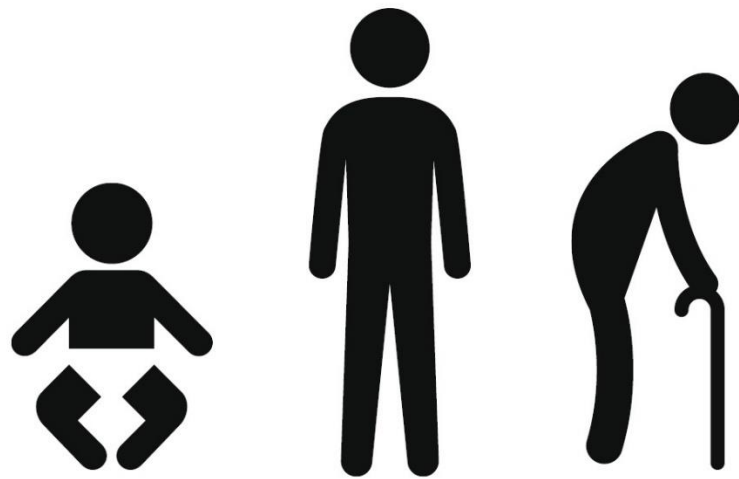
EDITORIAL AIR POLLUTION



CrossMark

The clear and persistent impact of air pollution on chronic respiratory diseases: a call for interventions

Isabella Annesi-Maesano¹, Francesco Forastiere², John Balmes^{3,4,5}, Erika Garcia⁶, Jack Harkema⁷, Stephen Holgate⁸, Frank Kelly², Haneen Khreis⁹, Barbara Hoffmann¹⁰, Cara Nichole Maesano¹, Rob McConnell¹¹, David Peden¹², Kent Pinkerton¹³, Tamara Schikowski¹⁴, George Thurston¹⁵, Laura S. Van Winkle¹⁶ and Christopher Carlsten¹⁷



Air pollution affects 100% of the population from unborn babies to the very elderly

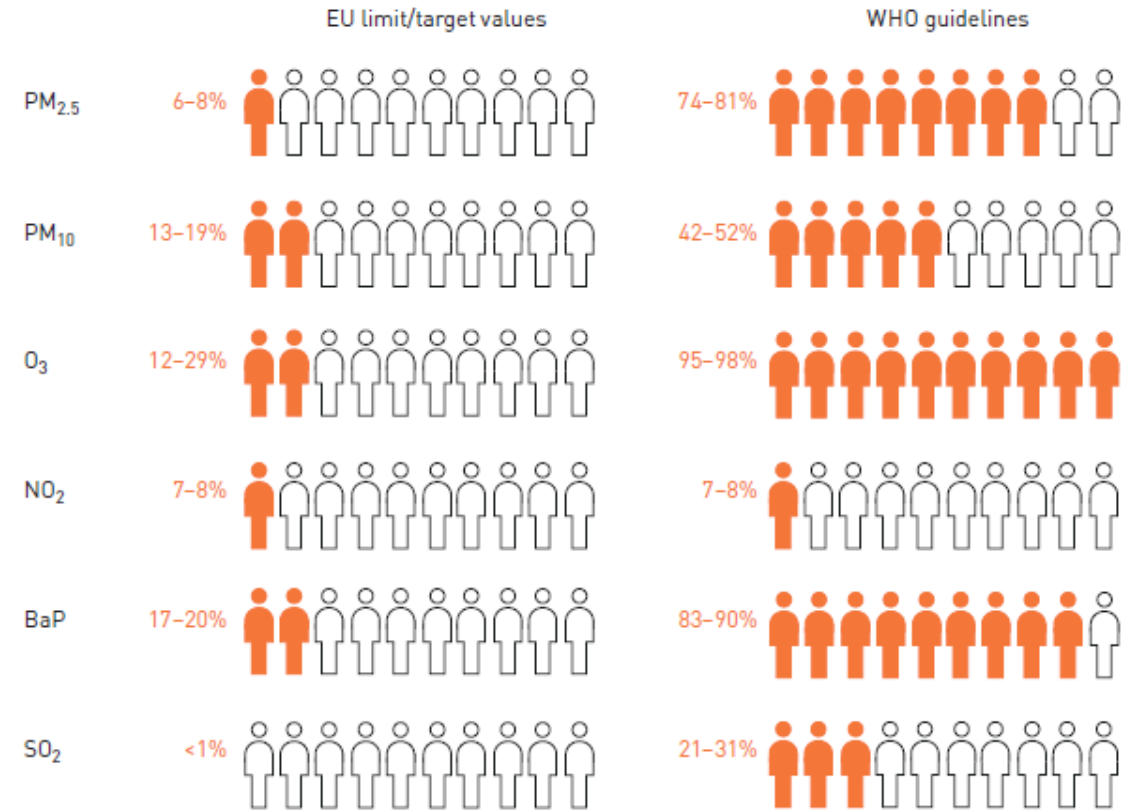
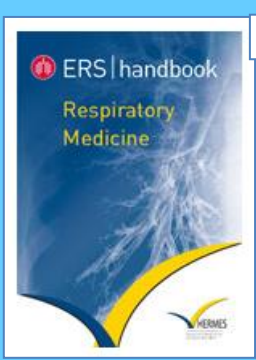


FIGURE 1 Exceedance of air quality standards and guidelines in European urban areas [data from www.eea.europa.eu/themes/air/health-impacts-of-air-pollution]. WHO: World Health Organization; PM_{2.5}: particulate matter of diameter of 2.5 µm, PM₁₀: particulate matter of diameter of 10 µm; BaP: benzo[a]pyrene. EU reference values (annual value): PM_{2.5}: 25 µg·m⁻³, PM₁₀: 40 µg·m⁻³, NO₂: 40 µg·m⁻³, O₃: 120 µg·m⁻³ (8-h mean); SO₂: 125 µg·m⁻³ (24-h mean); BaP: 1 ng·m⁻³. WHO air quality guidelines: EU reference values (annual value): PM_{2.5}: 10 µg·m⁻³, PM₁₀: 20 µg·m⁻³, NO₂: 40 µg·m⁻³, O₃: 100 µg·m⁻³ (8-h mean); SO₂: 20 µg·m⁻³ (24-h mean); BaP: 0.12 ng·m⁻³.



9/4/13

Indoor and outdoor pollution

Giovanni Viegi, Marzia Simoni, Sara Maio, Sonia Cerrai, Giuseppe Sarno and Sandra Baldacci

Table 1

Major outdoor/indoor pollutants and related health effects

Pollutant	Major sources	Health effects
Particulate matter	Outdoor Vehicular traffic Organic matter and fossil fuel combustion Power stations/industry Windblown dust from roadways, agriculture and construction Bushfires/dust storms	Lung cancer Premature death Mortality for cardiorespiratory diseases Reduced lung function Lower airway inflammation Upper airways irritation Neurological,
	Indoor Woodstoves Organic matter and fossil fuel combustion for heating/cooking ETS	cardiovascular diseases, metabolic disorders
Nitrogen dioxide	Outdoor Vehicular traffic Power stations/industry	Exacerbation of asthma Airway inflammation Bronchial hyperresponsiveness
	Indoor Unvented gas/kerosene appliances	Increased susceptibility to respiratory infection Reduced lung function

Ozone	Outdoor Sunlight chemical reaction with other pollutants Vehicular traffic Power stations/industry Consumer products	Lung tissue damage Reduced lung function Reduced exercise capacity Exacerbation of asthma Upper airway and eye irritation
	Carbon monoxide	Outdoor Organic matter and fossil fuel combustion Vehicular traffic Domestic heating Indoor Organic matter and fossil fuel combustion for heating/cooking Woodstoves Unvented gas/kerosene appliances ETS
Sulfur dioxide	Outdoor Coal/oil-burning power stations Industry/refineries Diesel engines Metal smelting	Exacerbation of respiratory diseases including asthma Respiratory tract irritation
VOCs	Indoor Building materials and products such as new furniture, solvents, paint, adhesives, insulation Cleaning activities and products Materials for offices	Lung cancer Asthma, dizziness, respiratory and lung diseases Chronic eye, lung or skin irritation Neurological and reproductive disorders

Viegi G, et al. Indoor and outdoor pollution. In "ERS Handbook, Respiratory Medicine 3rd Edition", Palange P, Rohde G eds., European Respiratory Society, Sheffield (UK) 2019: pp 771-778.

A joint ERS/ATS policy statement: what constitutes an adverse health effect of air pollution? An analytical framework

George D. Thurston¹, Howard Kipen², Isabella Annesi-Maesano³, John Balmes^{4,5}, Robert D. Brook⁶, Kevin Cromar⁷, Sara De Matteis⁸, Francesco Forastiere⁹, Bertil Forsberg¹⁰, Mark W. Frampton¹¹, Jonathan Grigg¹², Dick Heederik¹³, Frank J. Kelly¹⁴, Nino Kuenzli^{15,16}, Robert Laumbach², Annette Peters¹⁷, Sanjay T. Rajagopalan¹⁸, David Rich¹⁹, Beate Ritz²⁰, Jonathan M. Samet²¹, Thomas Sandstrom¹¹, Torben Sigsgaard²², Jordi Sunyer²³ and Bert Brunekreef^{13,24}

Eur Respir J 2017; 49: 1600419

- Newer evidences:
- Interstitial Pulmonary Fibrosis
- Exacerbations
- Incidence (?)
- Cystic fibrosis

FIGURE 1 Overview of diseases, conditions and biomarkers affected by outdoor air pollution. Updated based on [31]. Bold type indicates conditions currently included in the Global Burden of Disease categories.

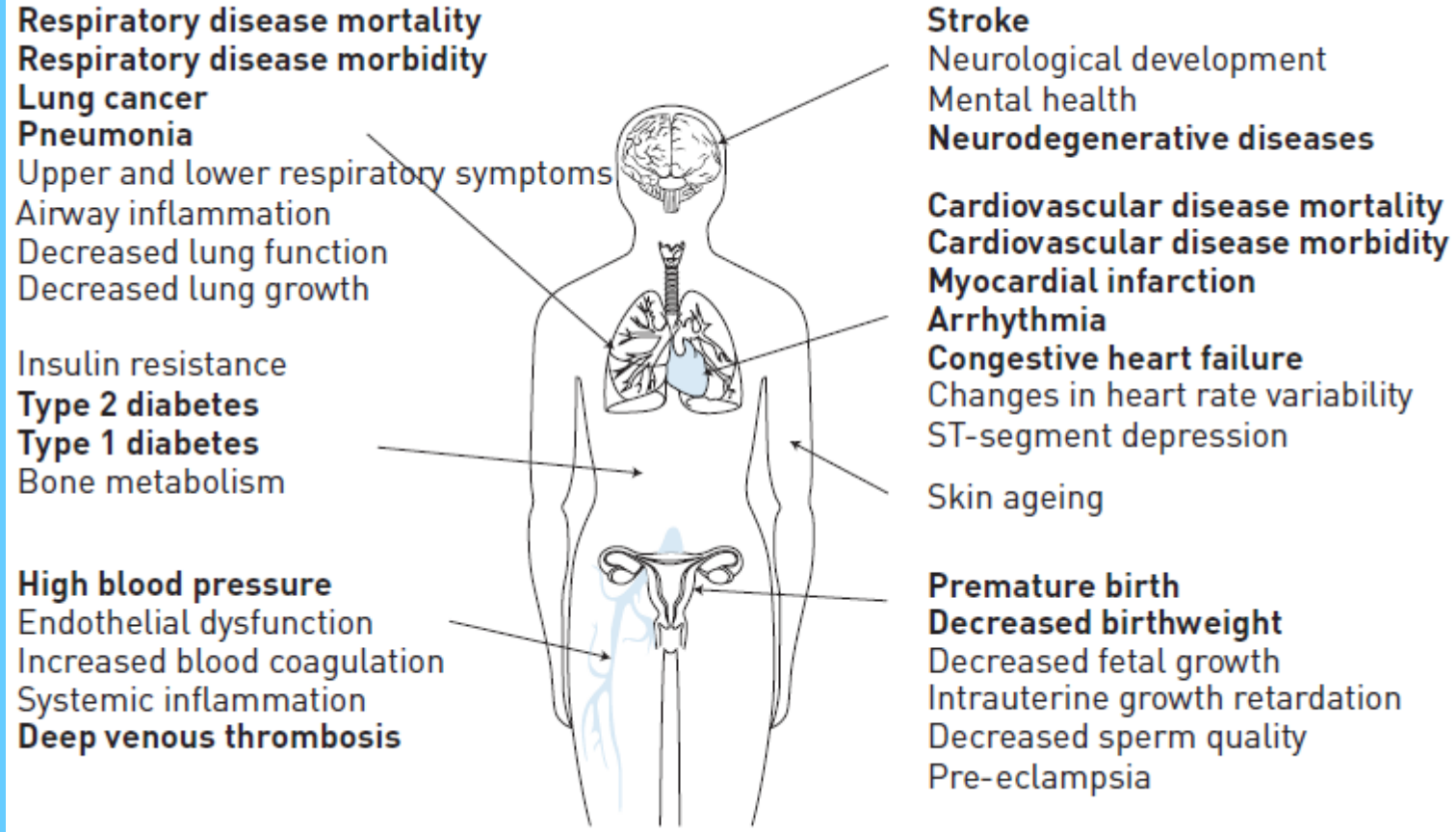


TABLE 3 Examples of respiratory clinical effects associated with air pollution

Increased respiratory mortality
Increased incidence of malignancies of the respiratory tract
Increased incidence, prevalence or frequency of exacerbations in chronic pulmonary disease: asthma, COPD and cystic fibrosis
Increased incidence or severity of upper and lower respiratory tract infections
Increased respiratory symptoms that affect quality of life: cough, phlegm, wheezing, dyspnoea and nasal drainage
Increased incidence of preterm birth, low birthweight or growth restriction leading to adverse respiratory outcomes
Reduced growth of lung function in children
Transient (hours) reductions in lung function associated with symptoms in healthy individuals
Transient (hours) reductions in lung function without symptoms in especially susceptible individuals (e.g. children with severe asthma)
Persistent or chronic (weeks, months or years) reductions in lung function

COPD: chronic obstructive pulmonary disease.

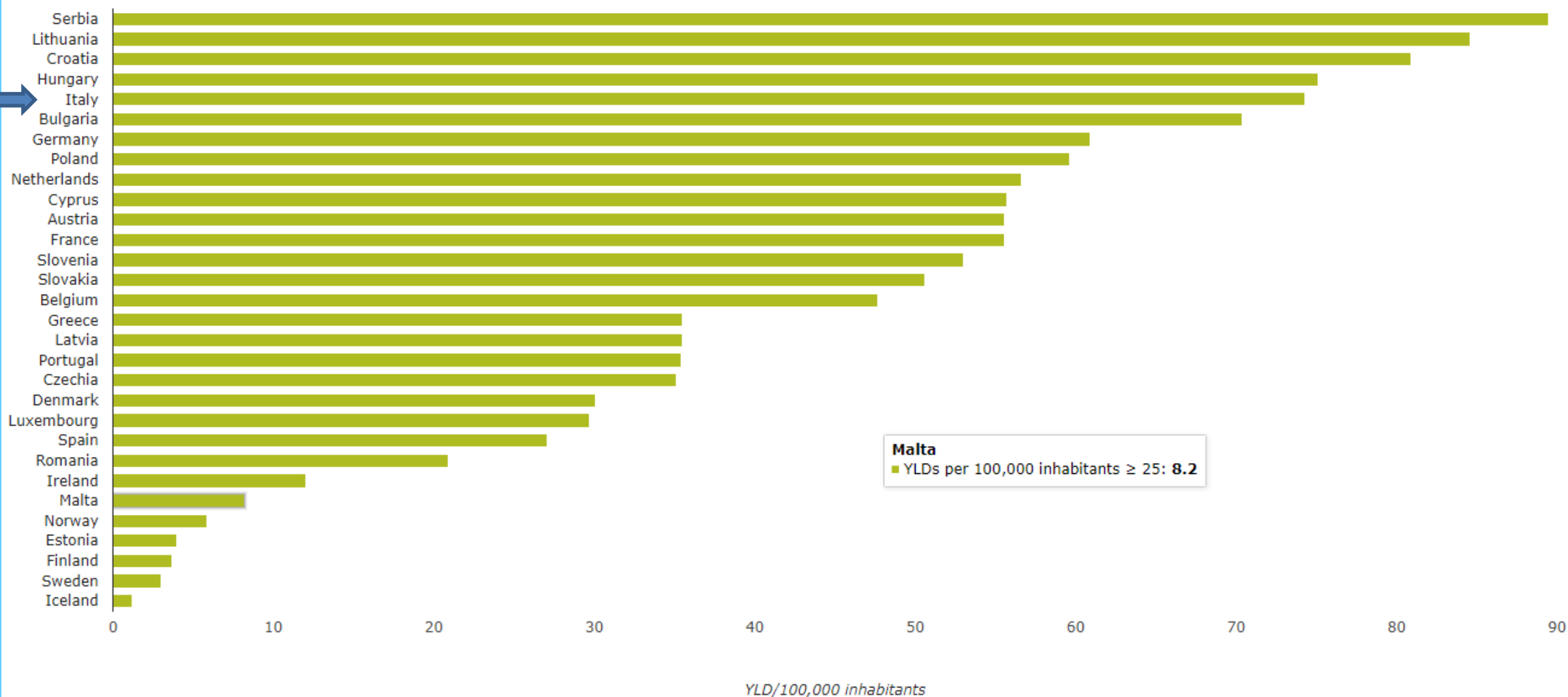
TABLE 4 Examples of biomarkers of potentially adverse respiratory health effects

Increased levels of markers of airway inflammation (e.g. PMNs or inflammatory cytokines in BAL or sputum)
Increased levels of markers of airway injury or inflammation in exhaled breath (e.g. increased acidity of exhaled breath condensate or increased F_{eNO} in asthmatics)
Increased levels of blood markers of lung injury (e.g. 8-isoprostanes, club cell secretory protein)
Imaging evidence for lung injury or reduced lung volume
Reduced pulmonary gas exchange (e.g. $DLCO$, $DLNO$, P_{aO_2} , pulse oximetry)
Increased airways responsiveness to nonspecific challenge
Increased airways hyperresponsiveness in asthmatic patients

PMN: polymorphonuclear leukocyte; BAL: bronchoalveolar lavage; F_{eNO} : exhaled nitric oxide fraction; $DLCO$: diffusing capacity of the lung for carbon monoxide; $DLNO$: diffusing capacity of the lung for nitric oxide; P_{aO_2} : arterial oxygen tension.

Figure 2. YLDs due to chronic obstructive pulmonary disease per 100,000 inhabitants attributable to PM_{2.5} for adults aged 25 and above for 30 European

Chart — YLDs due to chronic obstructive pulmonary per 100,000 inhabitants attributable to PM_{2.5} for adults aged 25 years and above for 30 European countries

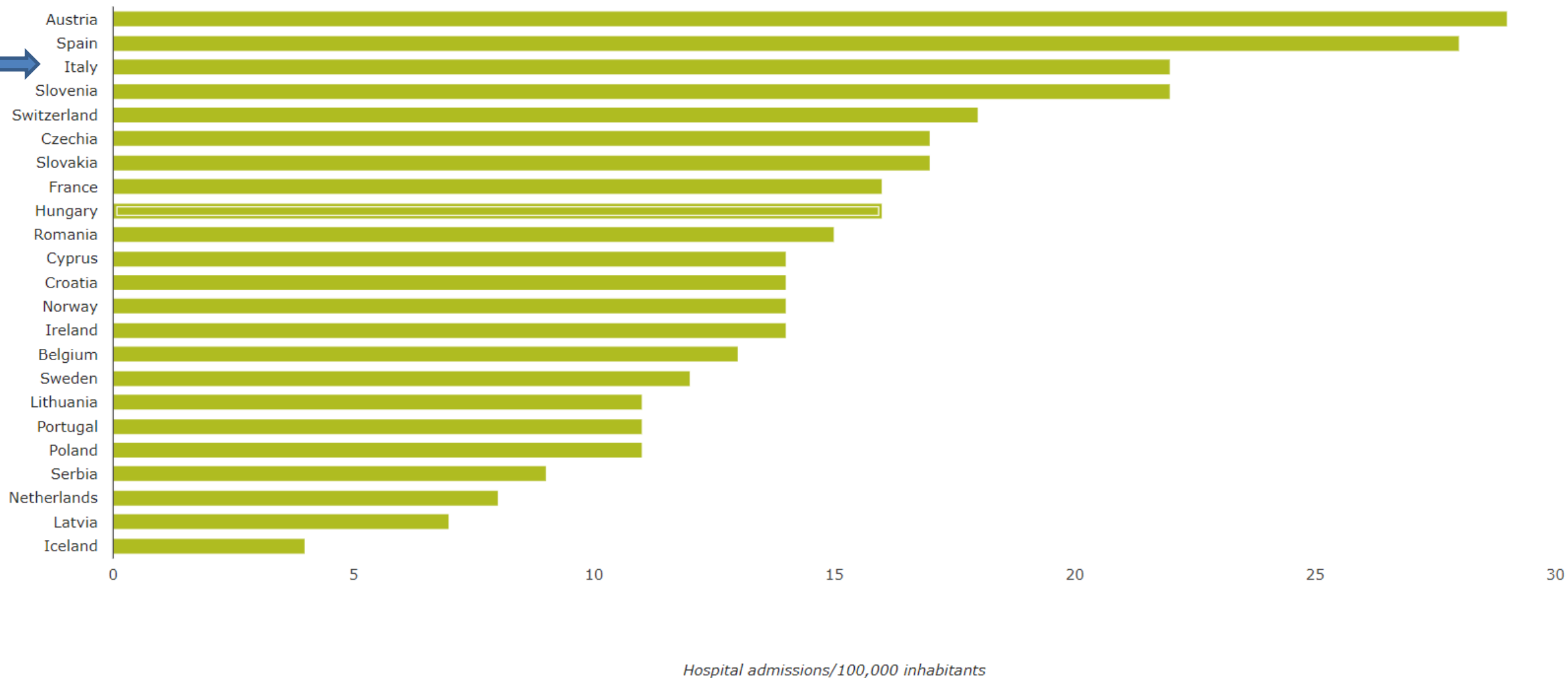


Note:

- YLDs: years lived with disability.
- Although 41 European countries have been considered, based on the data availability, there was only possible to estimate the COPD disease burden for 30.

Figure 4. Hospital admissions for respiratory disease by 100,000 inhabitants attributable to O₃ for adults aged 65 and above for 23 European countries

Chart – Hospital admissions for respiratory disease by 100,000 inhabitants attributable to O₃ for adults aged 65 years and above for 23 European countries



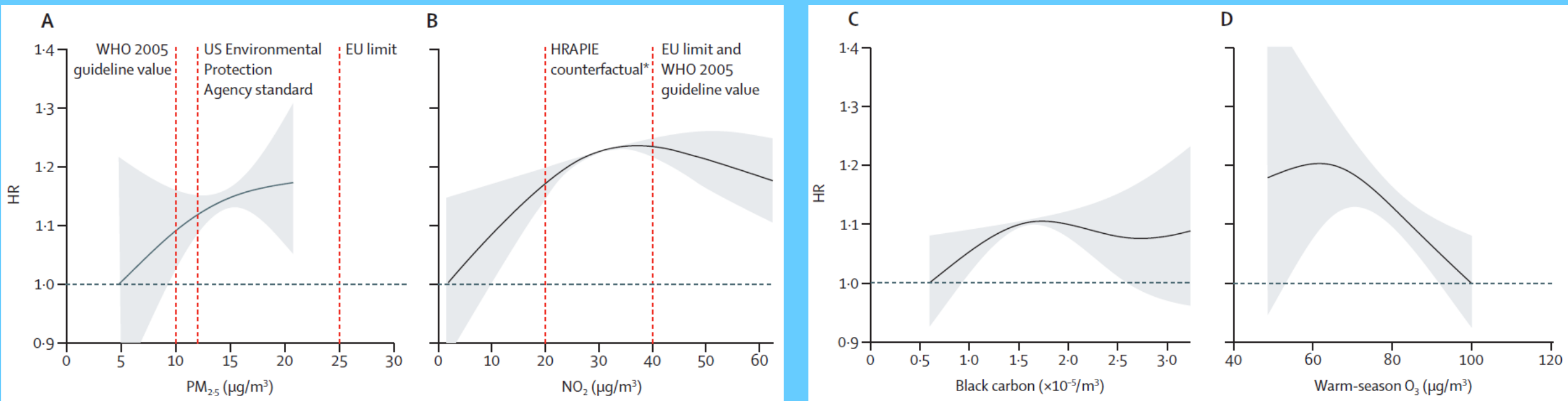
Note:

Although 41 European countries have been considered, based on the data availability, there was only possible to estimate the hospital admissions for 23.

Figure 3: Meta-analytical concentration-response functions of the association between air pollutants and non-accidental mortality

Cohort-specific models were adjusted for individual-level and area-level covariates available in the administrative cohorts (appendix p 11). Meta-analytical curves were obtained by meta-smoothing with natural splines with three degrees of freedom. The shaded regions are 95% CIs. HR=hazard ratio. NO₂=nitrogen dioxide. O₃=ozone. PM_{2.5}=fine particulate matter. *HRAPIE-suggested counterfactual below which no health impact is quantified.²¹

ELAPSE



New evidence: Economic and societal burden from air pollution is high

Joint Statement by Medical, Public Health and Scientific Societies

factors [8]. In terms of economic burden, the estimated global health-related external costs (i.e., those borne by society as a whole) were US\$ 5 trillion in 2013 with an additional US\$ 225 billion in lost labour productivity [9]. For the WHO European Region, the overall annual economic cost of health impacts and mortality from air pollution, including estimates for morbidity costs, stood at US\$ 1.575 trillion [10].

Costs from disease burden

Beneficial cost-benefit-ratio for clean air policies

Programmes that reduce air pollutant emissions provide enormous air quality and health benefits which increase over time. The estimated health benefits of improved air quality outweigh by far the implementation costs of air quality actions. For the US, it has been estimated that the benefits from decreased mortality, lower medical expenditures for air pollution-related diseases, and higher productivity of workers are around 30 times greater than the costs of the Clean Air Act, resulting in net improvements of economic growth and population welfare [19]. In China, public health benefits were

ITALIAN
EPIDEMIOLOGICAL
SURVEYS
(1980-2011)

CNR-IFC Study design: longitudinal, general population studies



PO Delta 1
(1980-82, n=3284, 8-64 yrs)
. Sampling
. CNR questionnaire
. Lung function test

Pisa 1
(1985-88, n=3865, 5-97 yrs)
. Sampling
. CNR questionnaire



PO Delta 2
(1988-91, n=2841, 8-73 yrs)
. CNR questionnaire
. Lung function test.
. Bronchial responsiveness
. Skin prick tests - Total serum IgE
. Nested: indoor

Pisa 2
(1991-93, n=2841, 8-97 yrs)
. CNR questionnaire
. Lung function tests
. Bronchial responsiveness
. Skin prick tests - Total serum IgE
. Mutagenetic determinations
. Nested indoor

SEASD*
(1997-98, n=2335, 13-99)
. Sub - sampling
. CNR questionnaire
. Blood sample collection
. Urine sample collection
. Blood pressure, height, weight

° Indicators
for Monitoring
COPD and
Asthma
in the EU

IMCAII°
(2006-11, n=1620, 18-103 yrs)
. Sub - sampling
. CNR questionnaire
. Lung function test
. Blood sample. Pulseoximeter
. Blood pressure, height, weight

General Population: Urban vs rural factor

Epidemiological Studies of Po Delta and Pisa

General Population: Urban vs sub-urban factor

The Proportional Venn Diagram of Obstructive Lung Disease in the Italian General Population* *Chest* 2004;126;1093-1101

*Giovanni Viegi, MD; Gabriella Matteelli, MD; Anna Angino, BS;
Antonio Scognamiglio, MD; Sandra Baldacci, BSc; Joan B. Soriano, MD, PhD;
and Laura Carrozzi, MD*

[CANCER RESEARCH 58, 4122-4126, September 15, 1998]

Serum Antibodies to Benzo(a)pyrene Diol Epoxide-DNA Adducts in the General Population: Effects of Air Pollution, Tobacco Smoking, and Family History of Lung Diseases¹

*Stefano Petruzzelli,² Alessandro Celi, Nolita Pulerà, Filomena Baliva, Giovanni Viegi, Laura Carrozzi,
Gigliola Ciacchini, Matteo Bottai, Francesco Di Pede, Paolo Paoletti, and Carlo Giuntini*

**Table 2—Prevalence Rates of CB, Emphysema, and
Asthma in the Two Italian General Population Samples**

Disease	Po River Delta, % (n = 2,463)	Pisa, % (n = 1,890)	p Value*
OLD	6.9	10.9	0.000
Asthma only	4.54	5.82	
Asthma + CB	0.28	0.21	
Asthma + emphysema	0.20	0.26	
CB only	0.89	1.22	
CB + emphysema	0.12	0.85	
Emphysema only	0.61	2.28	
CB + emphysema + asthma	0.24	0.21	

Zona rurale

Zona urbana

*By χ^2 test.

**Table 3 Multiple regression logistic analysis of the presence of serum anti-BPDE-DNA
antibodies with questionnaire variables**

	OR	95% CI
Urban residence	1.49	1.16-1.92
Tobacco smoking	1.25	1.06-1.48
Passive smoking	0.97	0.74-1.27
Family history of chronic bronchitis	1.02	0.65-1.60
Family history of emphysema	0.99	0.61-1.60
Family history of lung cancer	1.30	0.90-1.88
Other members of the family cluster with serum anti-BPDE-DNA antibodies	1.30	1.03-1.65

Urban residence is associated with bronchial hyper-responsiveness in Italian general population samples

Sara Maio, Sandra Baldacci, Laura Carrozzi, Eva Polverino, Anna Angino, Francesco Pistelli, Francesco Di Pede, Marzia Simoni, Duane Sherrill and Giovanni Viegi
Chest 2009;135;434-441

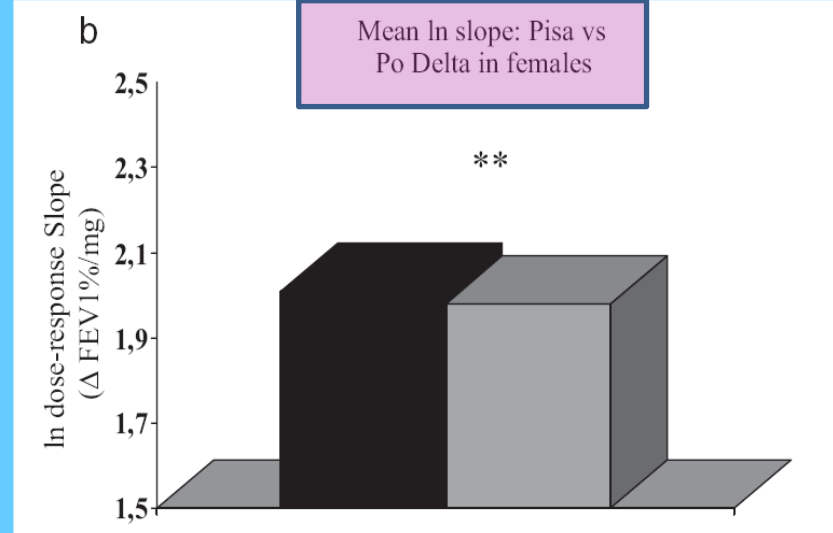
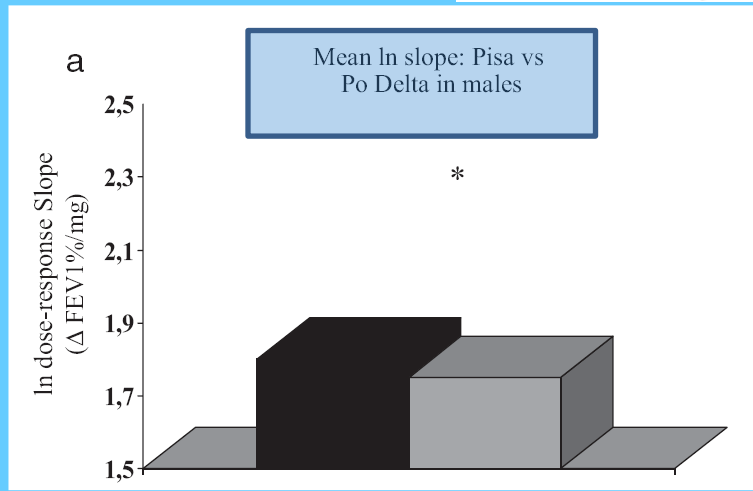


FIGURE 1. Top, a: * $p < 0.05$ with ANOVA. Bottom, b: ** $p < 0.01$ with ANOVA. Gray columns indicate Po Delta; black columns indicate Pisa.

Table 5. Effect of the independent variables on ln dose-response slope

	Odds Ratio	95% Confidence Interval
Gender:		
male	1	-
female	1.97	1.57-2.46
Groups of age:		
8-14	2.52	1.52-4.20
15-24	1.43	1.03-1.99
25-34	1	-
35-44	0.86	0.61-1.21
45-54	0.91	0.65-1.29
55-64	1.22	0.84-1.77
65-74	1.08	0.58-2.00
Smoking habits:		
never smoking	1	-
current smoking	1.39	1.05-1.83
ex smoking	1.11	0.84-1.46
Respiratory symptoms/diseases:		
others	1	-
chronic bronchitis	1.30	0.94-1.78
asthma	2.65	1.93-3.64
Prick test:		
negative	1	-
positive	1.32	1.05-1.67
log IgE values:		
< 1.93*	1	-
≥ 1.93	1.61	1.25-2.06
Residence:		
rural	1	-
urban	1.41	1.13-1.76
Airway caliber		
	0.66	0.61-0.73

*corresponding real number = 85.11 kU/l

Geographical information system and environmental epidemiology: a cross-sectional spatial analysis of the effects of traffic-related air pollution on population respiratory health

Daniela Nuvolone^{1,2*}, Roberto della Maggiore², Sara Maio³, Roberto Fresco², Sandra Baldacci³, Laura Carrozzi³, Francesco Pistelli³, Giovanni Viegi^{3,4} *Environmental Health* 2011, **10**:12

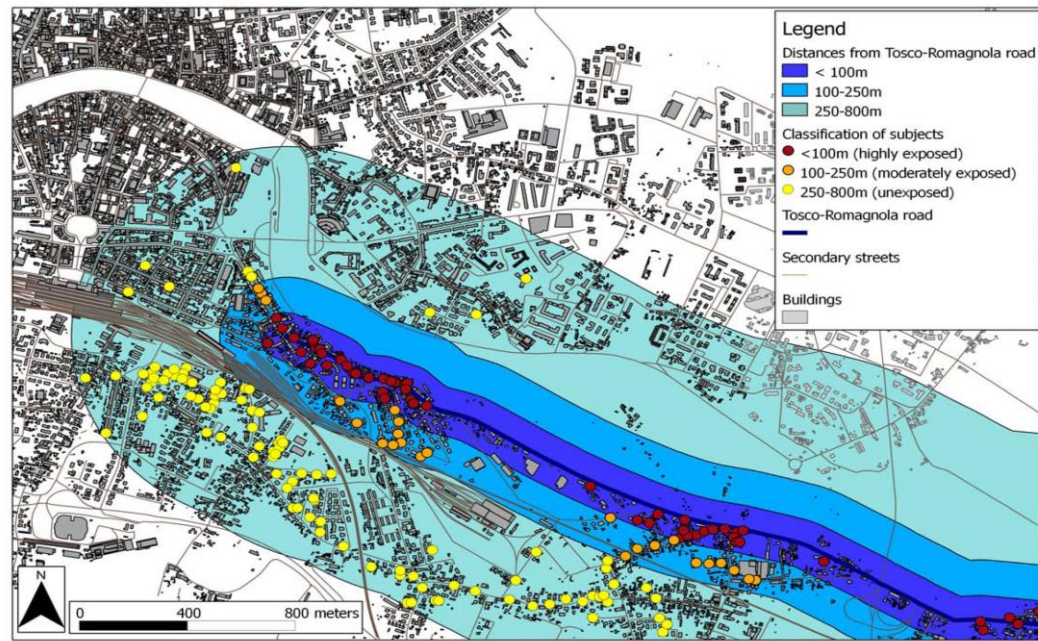


Figure 2 Classification of subjects based on the distance of each home from the main road. Zoomed map representing the classification of subjects according to the distance of each home from the main road. Highly exposed subjects are those living in the buffer area 0-100 m from the road, moderately exposed subjects living in the buffer area 100-250 m and unexposed are those living between 250 and 800 m from the road.

Table 6 Effects of distance of residence to main road on respiratory symptoms/diseases and dichotomized test outcomes: OR[†] and 95% CI

	Males		Females	
	<100 m	100-250 m	<100 m	100-250 m
<u>Persistent wheeze</u>	<u>1.76 * (1.08-2.87)</u>	1.54 # (0.94-2.53)	1.32 (0.76-2.28)	0.77 (0.42-1.42)
<u>Dyspnea</u>	0.88 (0.55-1.41)	0.86 (0.59-1.53)	<u>1.61 ** (1.13-2.27)</u>	1.35 # (0.95-1.93)
<u>COPD</u>	<u>1.80 * (1.03-3.08)</u>	1.21 (0.69-2.13)	1.60 (0.71-3.59)	0.99 (0.39-2.51)
Asthma	1.59 (0.85-2.98)	1.55 (0.83-2.87)	1.68 # (0.97-2.88)	0.58 (0.30-1.15)
Attacks of shortness of breath with wheeze	1.47 (0.87-2.48)	1.20 (0.70-2.04)	1.67 # (0.98-2.84)	0.74 (0.39-1.38)
<u>Skin test 5 mm pos.</u>	1.07 (0.67-1.72)	1.10 (0.70-1.73)	<u>1.83 * (1.11-3.00)</u>	0.95 (0.57-1.60)
<u>FEV₁/FVC% <70%</u>	<u>2.07 * (1.11-3.87)</u>	<u>2.53 ** (1.42-4.53)</u>	1.01 (0.48-2.14)	0.88 (0.41-1.89)
FEV ₁ /VC% <70%	1.15 (0.63-2.11)	1.76 * (1.02-3.04)	0.84 (0.40-1.72)	0.48 (0.21-1.11)

† OR adjusted for age, educational level, smoking habits, passive smoking exposure, occupational exposure, working position, number of hours spent at home and time of residence, calculated with subjects living between 250-800 m as the reference group.

*** p < 0.001, ** p < 0.01, * p < 0.05, # 0.05 < p < 0.1 (borderline).

Respiratory symptoms/diseases prevalence is still increasing: a 25-yr population study

Sara Maio ^{a, *}, Sandra Baldacci ^a, Laura Carrozzi ^b, Francesco Pistelli ^b, Anna Angino ^a, Marzia Simoni ^a, Giuseppe Sarno ^a, Sonia Cerrai ^a, Franca Martini ^a, Martina Fresta ^a, Patrizia Silvi ^a, Francesco Di Pede ^a, Massimo Guerriero ^c, Giovanni Viegi ^{a, d}

Respiratory Medicine 110 (2016) 58–65

Table 3
Risk factors for asthma/allergic rhinitis symptoms/diagnoses: OR and 95% CI.

	Asthma diagnosis	Attacks of asthma	Allergic rhinitis
<i>Survey:</i>			
PI1	1.00	1.00	1.00
PI2	1.08 (0.94–1.25)	0.88 (0.71–1.10)	1.26 (1.13–1.40)
PI3	1.34 (1.09–1.66)	1.90 (1.46–2.47)	2.98 (2.58–3.44)
Age	1.000 (0.991–1.001)	1.010 (1.003–1.020)	0.996 (0.992–0.999)
<i>Sex:</i>			
Females	1.00	1.00	1.00
Males	1.00 (0.80–1.26)	0.91 (0.69–1.21)	0.90 (0.78–1.04)
<i>Work exposure:</i>			
No	1.00	1.00	1.00
Yes	1.23 (1.03–1.46)	1.27 (1.01–1.60)	1.37 (1.22–1.55)
<i>Pack-years:</i>			
0	1.00	1.00	1.00
≤7	1.05 (0.82–1.36)	1.30 (0.92–1.85)	1.08 (0.92–1.28)
8–24	0.97 (0.73–1.27)	1.23 (0.86–1.74)	0.89 (0.75–1.06)
≥24	1.23 (0.92–1.64)	2.04 (1.47–2.84)	0.88 (0.73–1.07)
<i>Educational level:</i>			
>13 yrs	1.00	1.00	1.00
9–13 yrs	0.79 (0.52–1.19)	0.83 (0.47–1.47)	0.88 (0.69–1.13)
<8 yrs	1.12 (0.75–1.67)	1.28 (0.75–2.18)	0.75 (0.59–0.96)
<i>Area:</i>			
Suburban	1.00	1.00	1.00
Urban	0.89 (0.73–1.10)	1.10 (0.87–1.40)	1.19 (1.05–1.35)

PI1 = Pisa 1 survey; PI2 = Pisa 2 survey; PI3 = Pisa 3 survey.
OR and 95% CI from the multivariate generalised estimating equations.
Statistically significant values are represented in bold.

Table 4
Risk factors for COPD symptoms/diagnoses and airway obstruction^o: OR and 95% CI.

	Usual cough	Usual phlegm	COPD*	LLN airway obstruction [#]
<i>Survey:</i>				
PI1	1.00	1.00	1.00	
PI2	1.11 (0.98–1.25)	1.13 (0.99–1.29)	1.24 (1.02–1.52)	1.00
PI3	1.10 (0.93–1.30)	1.48 (1.25–1.75)	1.46 (1.14–1.85)	1.78 (1.40–2.27)
Age	1.015 (1.011–1.019)	1.019 (1.014–1.023)	1.050 (1.042–1.058)	1.022 (1.013–1.031)
<i>Sex:</i>				
Females	1.00	1.00	1.00	1.00
Males	0.94 (0.80–1.11)	1.36 (1.15–1.61)	1.55 (1.17–2.05)	0.76 (0.57–1.01)
<i>Work exposure:</i>				
No	1.00	1.00	1.00	1.00
Yes	1.25 (1.10–1.44)	1.40 (1.22–1.62)	1.81 (1.46–2.24)	1.22 (0.95–1.57)
<i>Pack-years:</i>				
0	1.00	1.00	1.00	1.00
≤7	1.85 (1.51–2.27)	1.80 (1.44–2.24)	1.26 (0.83–1.91)	1.81 (1.27–2.57)
8–23	2.66 (2.19–3.22)	2.67 (2.19–3.26)	2.25 (1.62–3.14)	2.16 (1.54–3.02)
≥24	4.44 (3.64–5.40)	4.64 (3.80–5.67)	4.45 (3.30–5.99)	2.69 (1.89–3.84)
<i>Educational level:</i>				
>13 yrs	1.00	1.00	1.00	1.00
9–13 yrs	1.25 (0.87–1.79)	0.97 (0.70–1.37)	1.24 (0.66–2.31)	0.94 (0.57–1.55)
≤8 yrs	1.57 (1.11–2.21)	1.11 (0.80–1.53)	1.39 (0.77–2.51)	1.06 (0.65–1.73)
<i>Area:</i>				
Suburban	1.00	1.00	1.00	1.00
Urban	1.14 (0.99–1.31)	1.30 (1.12–1.49)	1.54 (1.25–1.90)	0.86 (0.67–1.11)

PI1 = Pisa 1 survey; PI2 = Pisa 2 survey; PI3 = Pisa 3 survey.

^o airway obstruction values available in PI2 and PI3 surveys.

* diagnosis of COPD or emphysema or chronic bronchitis computed only in adult subjects.

[#] Lower Limit of Normal (LLN) according to American Thoracic Society (ATS)/European Respiratory Society (ERS) criterion [18]: forced expiratory volume in the first second (FEV₁)/forced vital capacity (FVC) < 5th percentile of the predicted value.

OR and 95% CI from the multivariate generalised estimating equations.

Statistically significant values are represented in bold. Borderline values are represented in italics.

18-yr cumulative incidence of respiratory/allergic symptoms/diseases and risk factors in the Pisa epidemiological study

Sara Maio ^{a,b,*}, Sandra Baldacci ^a, Laura Carrozzi ^c, Francesco Pistelli ^d, Marzia Simoni ^a, Anna Angino ^a, Stefania La Grutta ^e, Vito Muggeo ^b, Giovanni Viegi ^{a,e}

Table 4a
Longitudinal risk factors for asthma/allergic symptom/disease incidence: OR and 95% CI.

	Asthma diagnosis	Asthma attacks	Wheeze	Allergic rhinitis
Smoking habits:				
never	1.0	1.0	1.0	1.0
persistent	0.7 (0.2–3.0)	2.7 (1.1–6.4)	1.7 (0.6–4.7)	0.9 (0.5–1.6)
remittent for <18 years	1.1 (0.3–3.6)	1.5 (0.6–3.8)	0.2 (0.0–1.4)	1.1 (0.7–1.9)
remittent for ≥18 years	1.0 (0.4–2.7)	1.4 (0.7–3.1)	1.0 (0.4–2.6)	1.0 (0.7–1.6)
incident	–	0.9 (0.1–7.6)	0.8 (0.1–7.1)	0.7 (0.2–2.1)
Occupational exposure:				
never	1.0	1.0	1.0	1.0
persistent	4.4 (1.4–13.6)	1.1 (0.5–2.6)	0.5 (0.1–1.7)	1.8 (1.1–3.0)
remittent	–	0.8 (0.2–2.9)	0.3 (0.0–2.6)	0.7 (0.4–1.9)
incident	1.8 (0.7–4.8)	0.9 (0.5–1.9)	1.0 (0.4–2.4)	1.6 (1.1–2.4)
Vehicular traffic exposure:				
never	1.0	1.0	1.0	1.0
persistent	1.3 (0.3–5.1)	0.6 (0.2–1.6)	1.0 (0.3–2.9)	1.5 (0.9–2.5)
remittent	2.4 (0.5–10.2)	0.6 (0.2–2.2)	0.8 (0.2–3.9)	0.8 (0.4–1.6)
incident	2.6 (0.8–8.2)	2.2 (1.0–4.5)	1.5 (0.6–3.7)	1.8 (1.2–2.8)

A logistic regression model for each considered outcome was used to estimate the effect of longitudinal changes in risk factor exposure (smoking habits, occupational exposure and vehicular traffic exposure) on respiratory symptom/disease incidence, controlling for baseline factors closely related to the onset of respiratory symptom/disease (age, sex, body mass index -BMI, passive smoking, positivity to skin prick test, family history of allergic rhinitis and family history of respiratory diseases (asthma, chronic bronchitis or emphysema)).

Table 4b
Longitudinal risk factors for bronchitic symptom/disease incidence: OR and 95% CI.

	COPD	Usual phlegm	Usual cough	Dyspnoea	AO _{LLN}
Smoking habits:					
never	1.0	1.0	1.0	1.0	1.0
persistent	5.4 (2.3–12.5)	2.9 (1.7–5.1)	1.9 (1.0–3.5)	1.8 (1.1–3.0)	2.7 (1.0–7.4)
remittent for <18 years	3.3 (1.4–7.7)	0.8 (0.5–1.6)	1.0 (0.5–1.9)	1.5 (0.9–2.4)	1.1 (0.4–3.4)
remittent for ≥18 years	2.4 (1.2–5.1)	1.1 (0.7–1.7)	1.0 (0.6–1.7)	1.3 (0.9–2.0)	1.2 (0.5–2.8)
incident	–	0.8 (0.2–3.0)	1.7 (0.6–5.1)	0.9 (0.3–2.8)	–
Occupational exposure:					
never	1.0	1.0	1.0	1.0	1.0
persistent	1.9 (0.9–4.1)	1.8 (1.1–3.2)	1.4 (0.8–2.6)	1.3 (0.8–2.0)	2.0 (0.8–5.2)
remittent	–	0.4 (0.1–1.3)	0.4 (0.1–1.4)	0.8 (0.4–1.7)	1.3 (0.3–5.4)
incident	1.6 (0.9–3.0)	1.5 (1.0–2.4)	1.6 (1.0–2.5)	1.9 (1.3–2.8)	1.1 (0.5–2.6)
Vehicular traffic exposure:					
never	1.0	1.0	1.0	1.0	1.0
persistent	1.7 (0.7–3.9)	1.0 (0.6–1.7)	0.7 (0.4–1.3)	1.0 (0.6–1.6)	0.4 (0.2–1.1)
remittent	2.6 (0.9–7.0)	1.1 (0.6–2.2)	1.1 (0.6–2.2)	1.0 (0.6–1.9)	0.4 (0.1–1.8)
incident	2.4 (1.1–5.2)	1.3 (0.8–2.0)	0.9 (0.6–1.5)	1.2 (0.8–1.8)	0.5 (0.2–1.2)

A logistic regression model for each considered outcome was used to estimate the effect of longitudinal changes in risk factor exposure (smoking habits, occupational exposure and vehicular traffic exposure) on respiratory symptom/disease incidence, controlling for baseline factors closely related to the onset of respiratory symptom/disease (age, sex, body mass index -BMI, passive smoking, family history of allergic rhinitis and family history of respiratory diseases (asthma, chronic bronchitis or emphysema)).
COPD: Chronic Obstructive Pulmonary Disease; AO_{LLN}: Airway obstruction computed according to the lower limit of normal.
In italic: borderline values; in bold: statistically significant values.

Urban grey spaces are associated with increased allergy in the general population

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C N Maesano ⁵, I Annesi-Maesano ⁵, G Viegi ⁶

Table 4

Effects of 10% increase in residential exposure to urban greyness on allergic biomarkers/conditions and serum antibodies to BPDE-DNA adducts (n = 2070).

	OR (95% CI)
<i>Allergic biomarkers/conditions</i>	
SPT positivity	1.07 (1.02–1.13)
Reference category: negativity	1.00
<hr/>	
Perennial SPT positivity	1.05 (0.90–1.12)
Reference category: negativity	1.00
<hr/>	
Seasonal SPT positivity	1.12 (1.05–1.19)
Reference category: negativity	1.00
<hr/>	
Type of sensitization:	
polysensitization	1.11 (1.04–1.19)
monosensitization	1.03 (0.96–1.11)
Reference category: negativity	1.00
<hr/>	
Asthma/allergic rhinitis co-presence:	
asthma & allergic rhinitis	1.10 (0.98–1.23)
only allergic rhinitis	1.10 (1.04–1.17)
only asthma	1.07 (0.99–1.15)
Reference category: neither asthma nor allergic rhinitis	1.00
<hr/>	
SPT and asthma/allergic rhinitis co-presence:	
SPT positivity & asthma/allergic rhinitis	1.16 (1.00–1.25)
only SPT positivity	1.02 (0.95–1.09)
only asthma/allergic rhinitis	1.06 (1.00–1.12)
Reference category: neither SPT nor asthma/allergic rhinitis	1.00
<hr/>	
Log IgE value:	
≥1.81 kU/L	1.00 (0.95–1.05)
Reference category: < 1.81 kU/L	1.00
<hr/>	
<i>Exposure biomarker</i>	
Positivity to serum antibodies to BPDE-DNA adducts	1.07 (1.01–1.14)
Reference category: negativity	1.00



Findings

Per each **10%** increase in grey spaces coverage near home:



- + **7%** for SPT positivity
- + **7%** presence of BPDE-DNA
- + **10%** only allergic rhinitis
- + **11%** for polysensitization
- + **12%** for seasonal SPT positivity
- + **16%** co-presence of SPT positivity and asthma/allergic rhinitis

**Bando Ricerche in Collaborazione (BRiC)
Piano Attività di Ricerca 2016-2018**

The project



Coordinator:
Giovanni Viegi

June 29, 2017- December 28, 2019

Estimation of daily PM₁₀ and PM_{2.5} concentrations in Italy, 2013–2015, using a spatiotemporal land-use random-forest model

Massimo Stafoggia^{a,b,*}, Tom Bellander^b, Simone Bucci^a, Marina Davoli^a, Kees de Hoogh^{c,d}, Francesca de' Donato^a, Claudio Gariazzo^e, Alexei Lyapustin^f, Paola Michelozzi^a, Matteo Renzi^a, Matteo Scortichini^a, Alexandra Shtein^g, Giovanni Viegi^h, Itai Kloog^g, Joel Schwartzⁱ

Particulate matter (PM) air pollution is one of the major causes of death worldwide, with demonstrated adverse effects from both short-term and long-term exposure. Most of the epidemiological studies have been conducted in cities because of the lack of reliable spatiotemporal estimates of particles exposure in nonurban settings. The objective of this study is to estimate daily PM₁₀ (PM < 10 μm), fine (PM < 2.5 μm, PM_{2.5}) and coarse particles (PM between 2.5 and 10 μm, PM_{2.5-10}) at 1-km² grid for 2013–2015 using a machine learning approach, the Random Forest (RF). Separate RF models were defined to: predict PM_{2.5} and PM_{2.5-10} concentrations in monitors where only PM₁₀ data were available (stage 1); impute missing satellite Aerosol Optical Depth (AOD) data using estimates from atmospheric *ensemble* models (stage 2); establish a relationship between measured PM and satellite, land use and meteorological parameters (stage 3); predict stage 3 model over each 1-km² grid cell of Italy (stage 4); and improve stage 3 predictions by using small-scale predictors computed at the monitor locations or within a small buffer (stage 5). Our models were able to capture most of PM variability, with mean cross-validation (CV) R² of 0.75 and 0.80 (stage 3) and 0.84 and 0.86 (stage 5) for PM₁₀ and PM_{2.5}, respectively. Model fitting was less optimal for PM_{2.5-10}, in summer months and in southern Italy. Finally, predictions were equally good in capturing annual and daily PM variability, therefore they can be used as reliable exposure estimates for investigating long-term and short-term health effects.

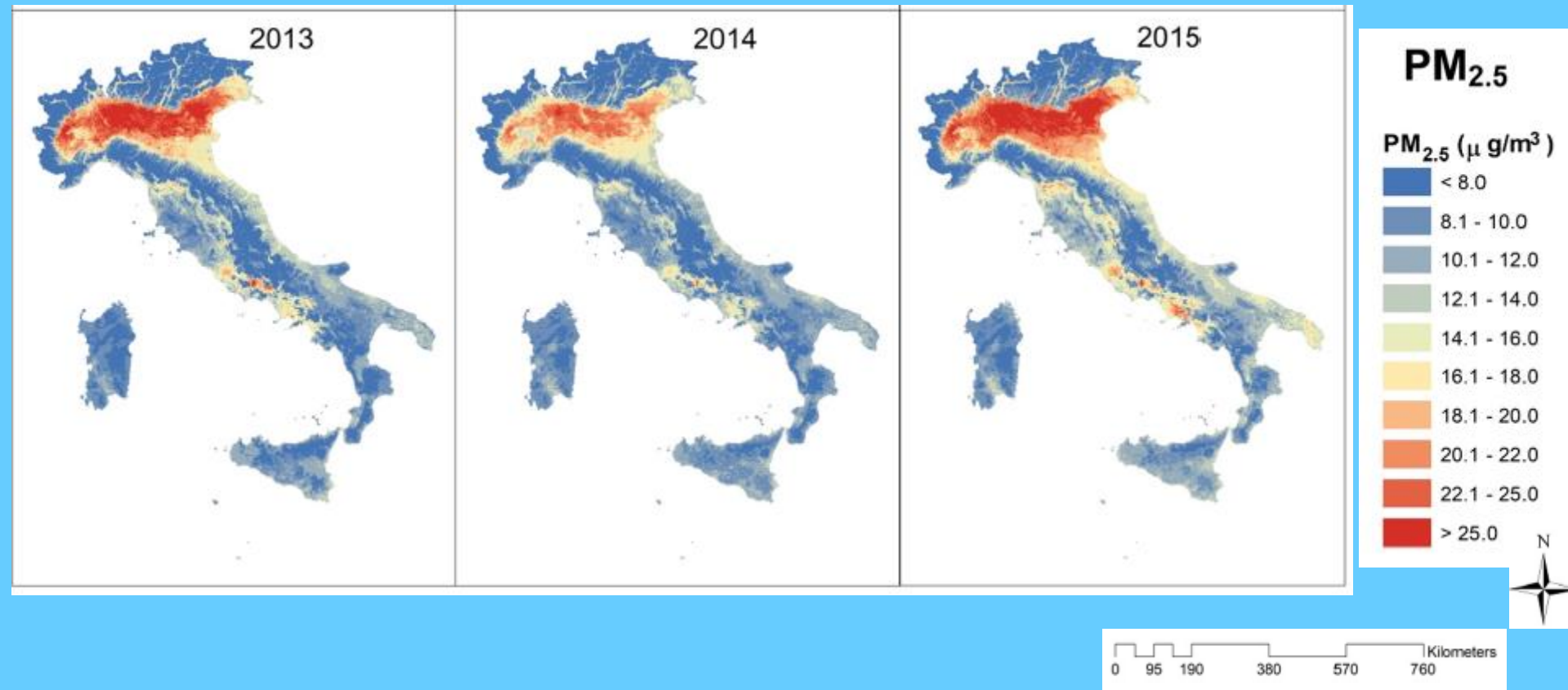


Fig. 2. Predicted PM_{10} (top) and $\text{PM}_{2.5}$ (bottom) concentrations from stage 4 model: annual means, 2013–2015.

Effects of Particulate Matter on the Incidence of Respiratory Diseases in the Pisan Longitudinal Study



Salvatore Fasola ^{1,*} , Sara Maio ², Sandra Baldacci ², Stefania La Grutta ¹, Giuliana Ferrante ³, Francesco Forastiere ¹, Massimo Stafoggia ⁴, Claudio Gariazzo ⁵ , Giovanni Viegi ^{1,2} and on behalf of the BEEP Collaborative Group [†]

Table 2. Associations (odds ratio, OR, and 95% confidence intervals (CI)) between risk factors ascertained during the first survey (1991–1993) and cumulative incidences of asthma, rhinitis, Chronic Obstructive Pulmonary Disease (COPD) and chronic phlegm ascertained at the second survey (2009–2011), from multivariable logistic regression models with Firth’s correction.

	Asthma	Rhinitis	COPD	Chronic Phlegm
Cumulative incidence:	4/284 (1.4%)	90/264 (34.1%)	29/282 (10.3%)	16/262 (6.1%)
Independent variables:	OR (95% CI)	OR (95% CI)	OR (95% CI)	OR (95% CI)
PM ₁₀ (1 µg/m ³ increase) ¹	₋₂	₋₂	2.96 (1.50–7.15)	₋₂
PM _{2.5} (1 µg/m ³ increase) ¹	₋₂	2.25 (1.07–4.98)	₋₂	4.17 (1.12–18.71)
Age, years (10-year increase)	₋₂	₋₂	1.87 (1.29–3.02)	₋₂
Male gender	₋₂	₋₂	₋₂	₋₂
Smoker (ref = non-smoker)	12.96 (1.25–∞)	₋₂	2.99 (1.08–9.39)	₋₂
Ex-smoker (ref = non-smoker)	4.86 (0.27–∞)	₋₂	1.67 (0.60–4.89)	₋₂
Occupational exposure	₋₂	₋₂	1.91 (0.83–4.79)	5.41 (1.88–21.79)

¹ Estimated exposure levels at the residential address for the year 2011, 1 km² resolution. ² Variables excluded by the stepwise selection procedure. Significant odds ratios are reported in bold.

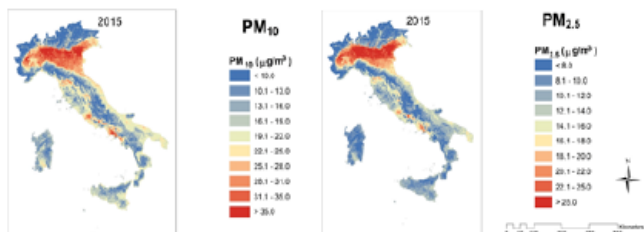
A nationwide study of air pollution from particulate matter and daily hospitalizations for respiratory diseases in Italy

Matteo Renzi ^{a,*,}, Matteo Scortichini ^{a,}, Francesco Forastiere ^{b, c,}, Francesca de' Donato ^{a,}, Paola Michelozzi ^{a,}, Marina Davoli ^{a,}, Claudio Gariazzo ^{c,}, Giovanni Viegi ^{b, d,}, Massimo Stafoggia ^{a,}, BEEP collaborative Group, Carla Ancona ^{f,}, Simone Bucchi ^{e,}, Francesca de' Donato ^{f,}, Paola Michelozzi ^{f,}, Matteo Renzi ^{f,}, Matteo Scortichini ^{f,}, Massimo Stafoggia ^{f,}, Michela Bonafede ^g ... Giuseppe Carlino ^h

Graphical abstract

A nationwide study of air pollution and daily hospitalizations for respiratory diseases in Italy

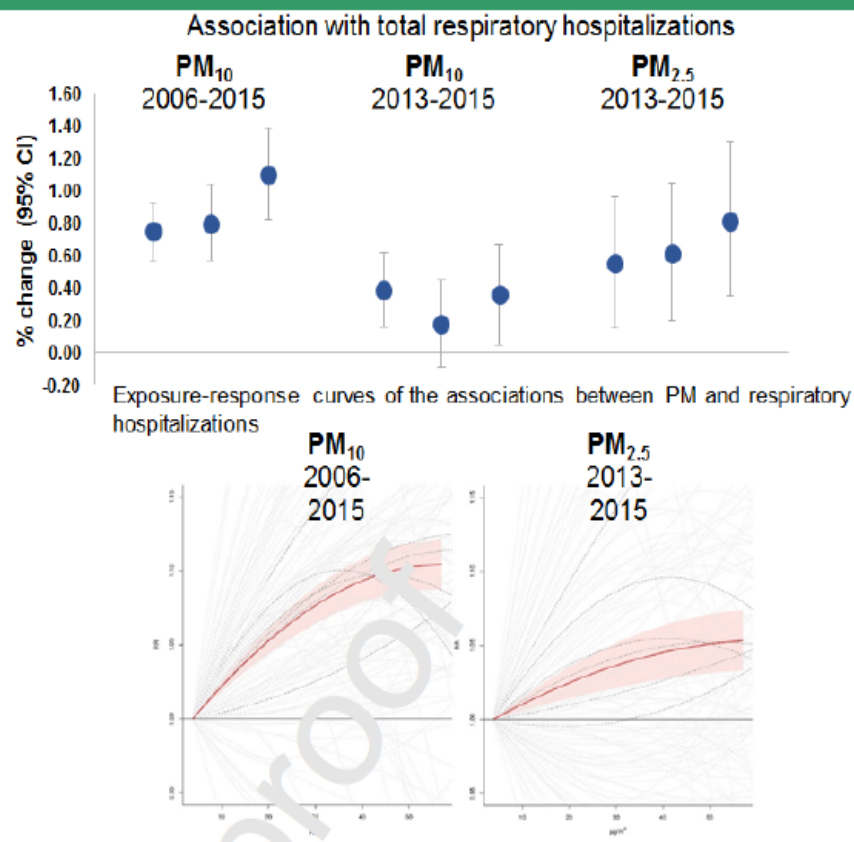
- Machine-learning approach to estimate PM exposure
- National health data database
- Sensitivity analyses (subgroup outcomes, effect modification, exposure-response curves)



Annual concentrations of PM10 and PM2.5 in Italy

Conclusions:

In this study we provided evidence of harmful effect of PM₁₀ and PM_{2.5} on respiratory hospitalizations in Italy during 2006–2015 and we reported a positive association for a subgroup of respiratory outcomes such as asthma, COPD and LRTI. Low-level effects were detected.



Abstract

Results

A total of 4,154,887 respiratory admission were registered during 2006–2015, of which 29% for LRTI, 12% for COPD, 6% for URTI, and 3% for asthma. Daily mean PM₁₀ and PM_{2.5} concentrations over the study period were 23.3 and 17 $\mu\text{g}/\text{m}^3$, respectively. For each 10 $\mu\text{g}/\text{m}^3$ increases in PM₁₀ and PM_{2.5} at lag 0–5 days, we found excess risks of total respiratory diseases equal to 1.20% (95% confidence intervals, 0.92, 1.49) and 1.22% (0.76, 1.68), respectively. The effects for the specific diseases were similar, with the strongest ones for asthma and COPD. Higher effects were found in the elderly and in less urbanized areas.

Conclusions

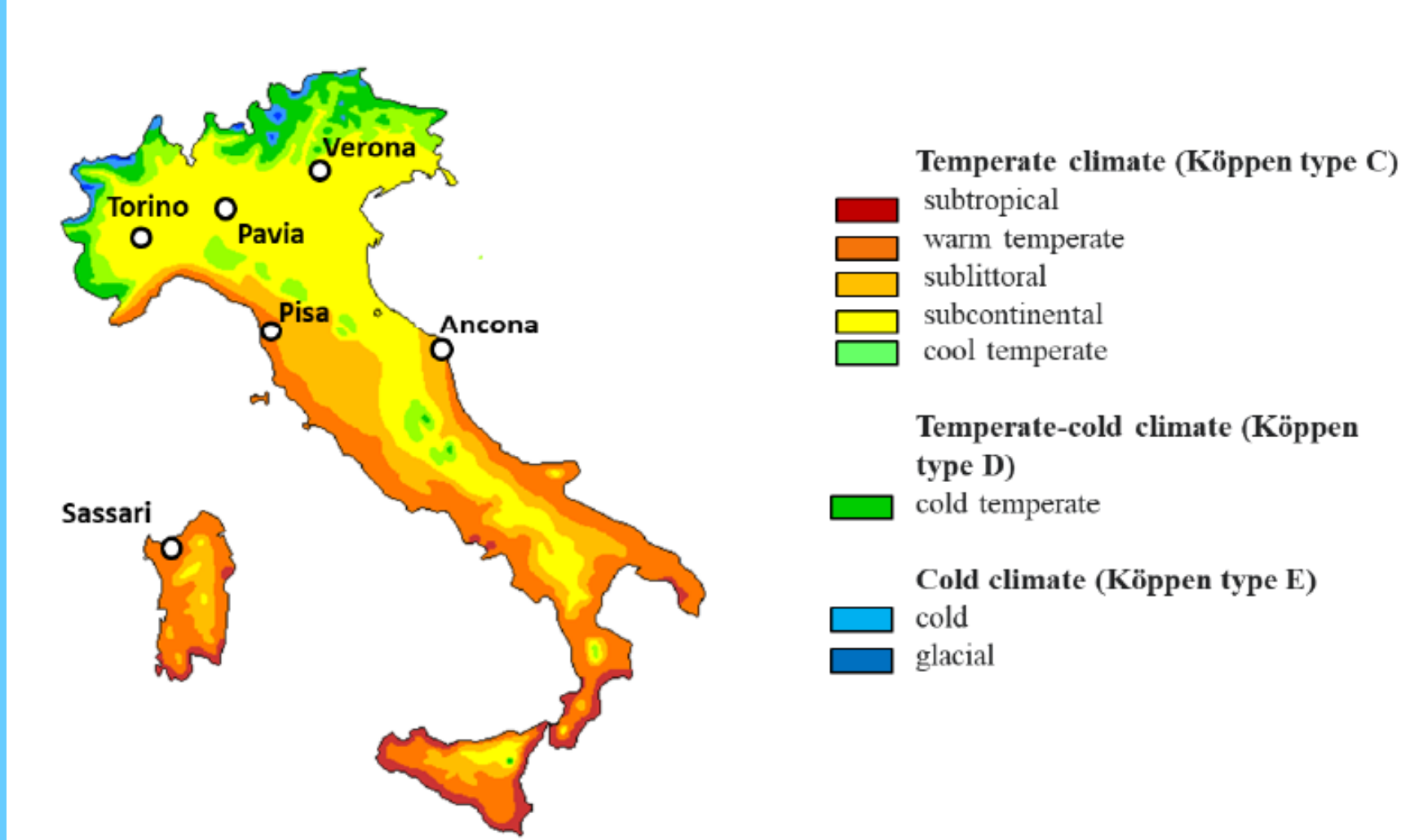
Short-term exposure to PM is harmful for the respiratory system throughout an entire country, especially in elderly patients. Strong effects can be found also in less urbanized areas.

Relationship of long-term air pollution exposure with asthma and rhinitis in Italy: an innovative multipollutant approach

Environmental Research 224 (2023) 115455

Sara Maio¹, Salvatore Fasola², Alessandro Marcon³, Anna Angino¹, Sandra Baldacci¹, Maria Beatrice Bilò⁴, Roberto Bono⁵, Stefania La Grutta², Pierpaolo Marchetti³, Giuseppe Sarno¹, Giulia Squillacioti⁵, Ilaria Stanisci¹, Pietro Pirina⁶, Sofia Tagliaferro¹, Giuseppe Verlati³, Simona Villani⁷, Claudio Gariazzo⁸, Massimo Stafoggia⁹, Giovanni Viegi¹, on behalf of the BIGEPI group*

Figure 1 Geographical distribution of the 6 cities



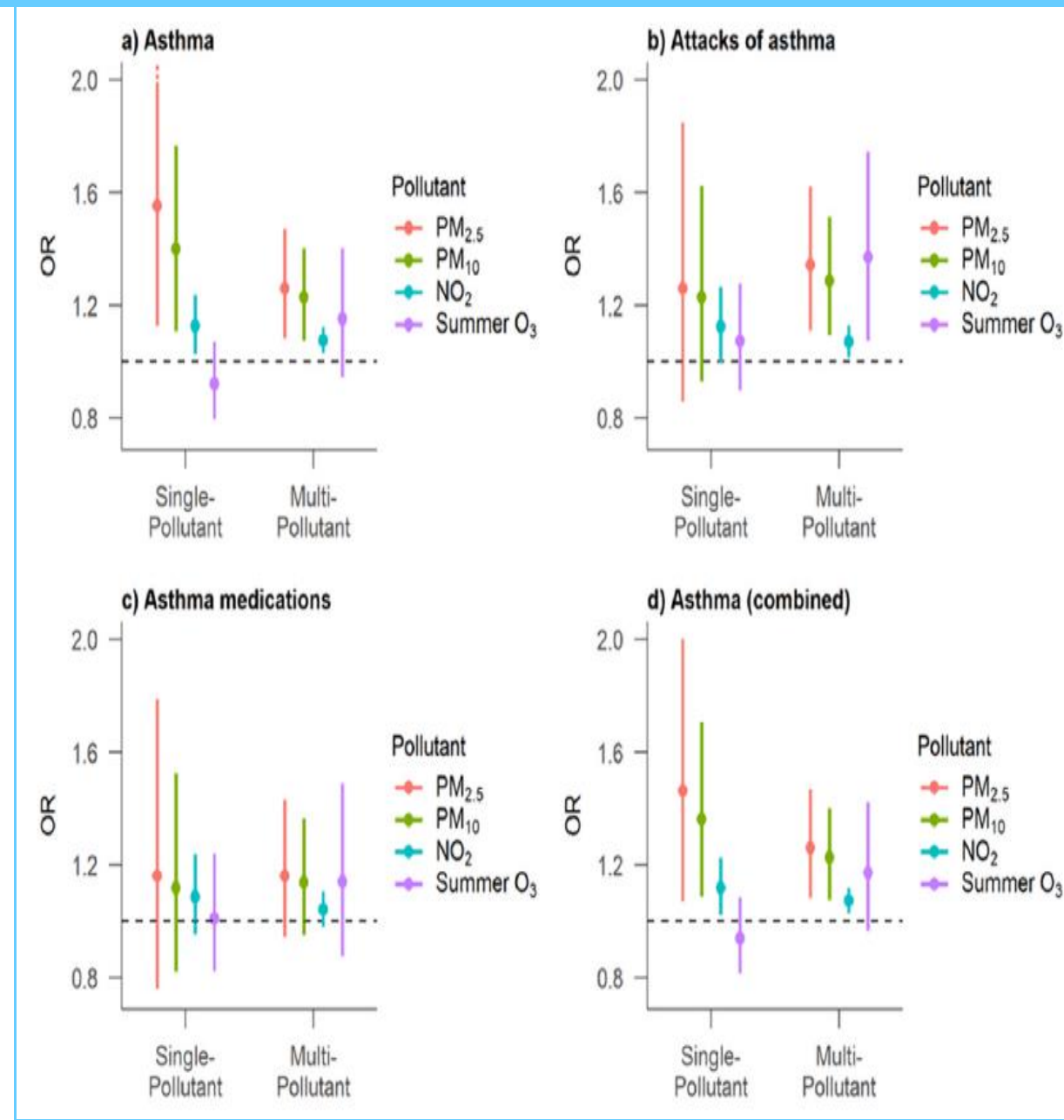
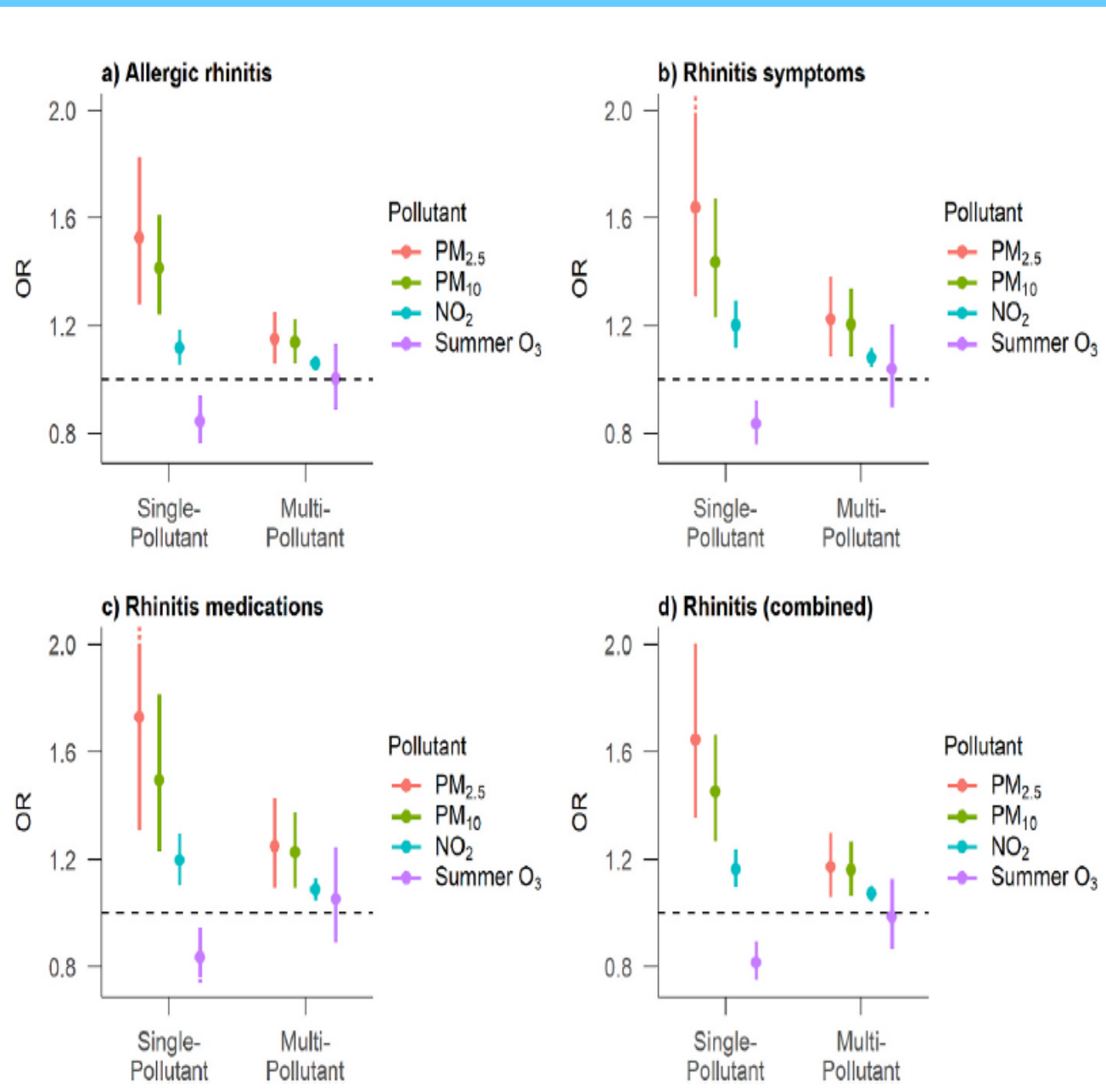


Fig. 3. Results of single-pollutant and multipollutant logistic regression models (OR and 95% CI for 10 µg/m³ increases): rhinitis.

GARD ITALIA

Sottogruppo di lavoro "Ambiente, Clima e Salute"

Ministero della Salute

Direzione Generale della Prevenzione Sanitaria

https://www.salute.gov.it/imgs/C_17_pubblicazioni_2945_allegato.pdf

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**NUOVE EVIDENZE A
SUPPORTO DEL DOCUMENTO
“INQUINAMENTO
ATMOSFERICO
E CAMBIAMENTI CLIMATICI
Elementi per una strategia
nazionale di prevenzione”
Aggiornamento 2023**

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PREMESSA

Nonostante il documento “INQUINAMENTO ATMOSFERICO E CAMBIAMENTI CLIMATICI” - Elementi per una strategia nazionale di prevenzione”, consultabile sul Sito del Ministero della Salute¹ sia relativamente recente (2019), il Comitato Esecutivo dell’Alleanza Globale contro le Malattie Croniche Respiratorie (GARD-Italia) ha sentito l’esigenza di aggiornarlo.

Le motivazioni sono da rinvenire negli importanti avvenimenti successivi al 2019, quali: la pandemia da COVID-19; la pubblicazione delle nuove linee guida per la qualità dell’aria emanate dall’Organizzazione Mondiale della Sanità (WHO); la serie di pubblicazioni Lancet Countdown; i nuovi documenti dell’*Intergovernmental Panel on Climate Change* (IPCC); il programma *Next Generation* EU; il Piano Nazionale di Ripresa e Resilienza (PNRR); la realizzazione di nuove ricerche scientifiche e la pubblicazione dei loro risultati.

Il sottogruppo di lavoro GARD-Italia che ha curato l’aggiornamento risulta così composto:

- **Fabrizio Anatra** - Ministero della Salute -Direzione Generale della Prevenzione Sanitaria
- **Gennaro D’Amato** - AIPO-ITS
- **Francesca De Maio** - ISPRA -Istituto Superiore Protezione e Ricerca Ambientale
- **Daniela Galeone** - Ministero della Salute -Direzione Generale della Prevenzione Sanitaria
- **Paolo Lauriola** - ISDE
- **Giovanni Viegi** - CNR e GARD Internazionale

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Si ringrazia il Gruppo *Minds for one health* (M4OH), di cui sono stati ripresi alcuni documenti pubblicati nel 2022-23.

Si ringraziano anche i componenti del Comitato Esecutivo GARD-Italia per gli utili commenti.

Tabella 1. Confronto tra nuove Linee guida OMS, precedenti Linee guida OMS e limiti normativi

		OMS – Linee guida per la qualità dell’aria 2005	OMS – Linee guida per la qualità dell’aria 2021	Direttive europee per la qualità dell’aria – Valori limite
PM _{2,5}	Annuali	10 µg/m ³	5 µg/m ³	25 µg/m ³
PM _{2,5}	Giornaliere (24 ore)	25 µg/m ³	15 µg/m ³	-
PM ₁₀	Annuali	20 µg/m ³	15 µg/m ³	40 µg/m ³
PM ₁₀	Giornaliere (24 ore)	50 µg/m ³	45 µg/m ³	50 µg/m ³
NO ₂	Annuali	40 µg/m ³	10 µg/m ³	40 µg/m ³
NO ₂	Giornaliere (24 ore)	-	25 µg/m ³	50 µg/m ³

Tabella 2. Morti premature attribuibili ad esposizione a PM_{2,5}, NO₂ e ozono in Italia, 2019

PM _{2,5}		NO ₂		O ₃		
Popolazione (1.000)	Media annuale	Morti premature	Media annuale	Morti premature	SOMO35	Morti premature
59.817	14,5	49.900	20	10.640	6.657	3.170

Tabella 3. Anni di vita persi (YLL) attribuibili ad esposizione a PM_{2,5}, NO₂ e ozono in Italia, 2019

Paese	YLL	YLL/100.000 abitanti	YLL	YLL/100.000 abitanti	YLL	YLL/100.000 abitanti
Italia	504.500	843	107.600	180	33.200	55

Tabella 4. Benefici teorici sulla salute, in termini di riduzione delle morti premature, che sarebbero stati ottenuti nel 2019 nei 27 paesi europei se fossero stati soddisfatti i differenti valori limiti europei e le linee guida OMS per il PM_{2,5}.

	Morti premature dovute al PM_{2,5}	Riduzione delle morti premature in base ai livelli del 2019	% riduzione delle morti premature in base ai livelli del 2019	% riduzione delle morti premature in base ai livelli del 2005
Concentrazioni nel 2019	306.700	-	-	33%
Valori limite europei 25 µg/m³	306.500	200	0%	33%
Valori limite indicativi europei 20 µg/m³	303.500	3.200	1%	33%
Target intermedio 3 del 2021 OMS 15 µg/m³	289.200	17.500	6%	37%
Target intermedio 4 del 2021 OMS (linee guida sulla qualità dell'aria OMS 2005) 10 µg/m³	241.400	65.300	21%	47%
Linee guida sulla qualità dell'aria OMS 2021 5 µg/m³	129.400	177.300	58%	72%

NB: Il target intermedio e le linee guida OMS sono stati estrapolati dall'aggiornamento delle linee guida pubblicate nel 2021.

Tabella 5. Benefici teorici sulla salute, in termini di riduzione delle morti premature, che sarebbero stati ottenuti nel 2019 se fossero stati soddisfatti i differenti valori limite europei e le linee guida OMS per il PM_{2,5}, in Italia.

Livelli 2019	Valore limite europeo 25 µg/m³		Valore indicativo europeo 20 µg/m³		Target intermedio 3 OMS 15 µg/m³		Target intermedio 4 OMS (linee guida sulla qualità dell'aria OMS 2005) 10 µg/m³		Linee guida sulla qualità dell'aria OMS 5 µg/m³	
Morti premature	Morti premature	% riduzione dei livelli 2019	Morti premature	% riduzione dei livelli 2019	Morti premature	% riduzione dei livelli 2019	Morti premature	% riduzione dei livelli 2019	Morti premature	% riduzione dei livelli 2019
49.900	49.900	0	49.100	2	45.200	9	34.500	31	17.700	65

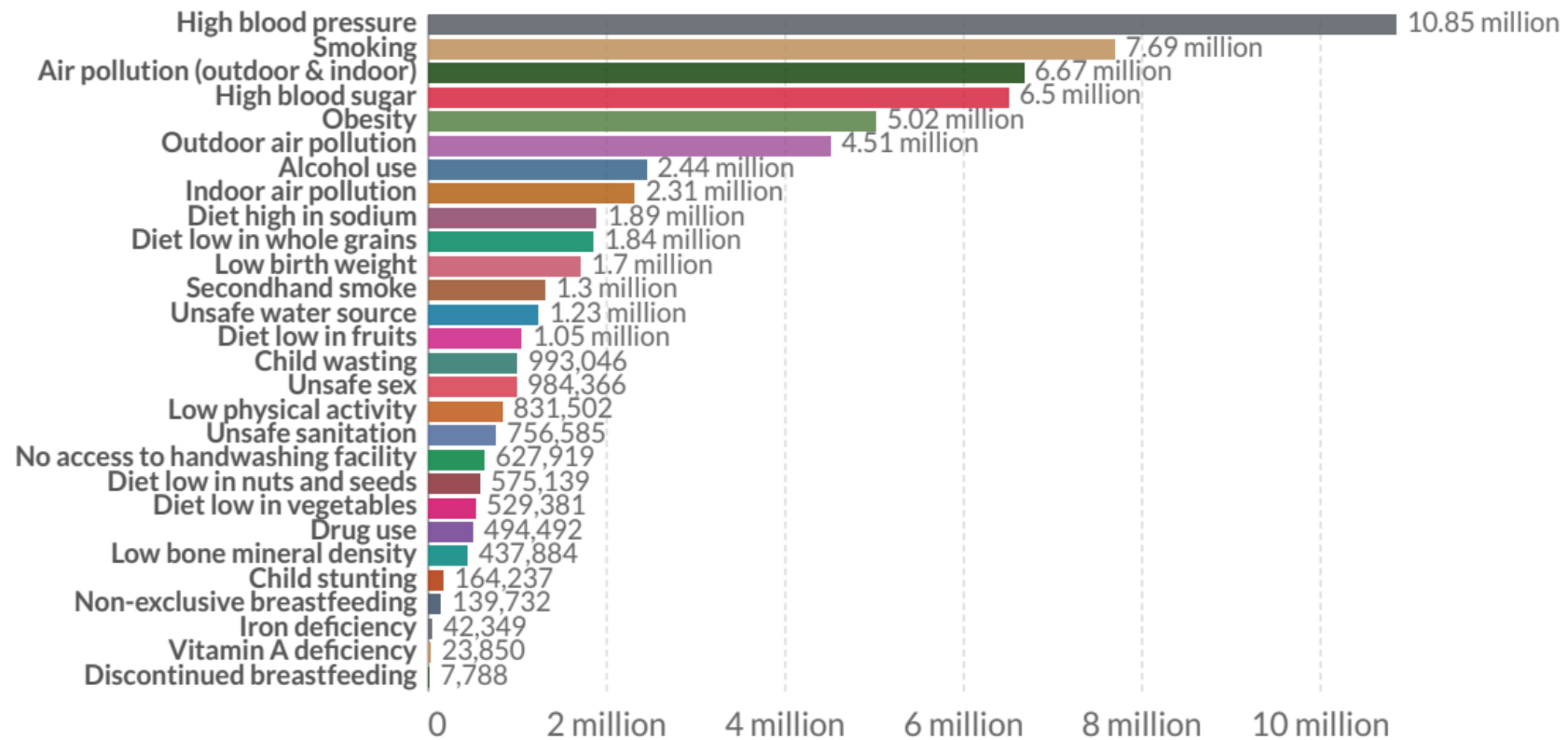
Figura 1. Numero di decessi per fattore di rischio, nel mondo 2019

Number of deaths by risk factor, World, 2019

Total annual number of deaths by risk factor, measured across all age groups and both sexes.

Our World
in Data

↔ Change country



Source: IHME, Global Burden of Disease (GBD)

OurWorldInData.org/causes-of-death • CC BY

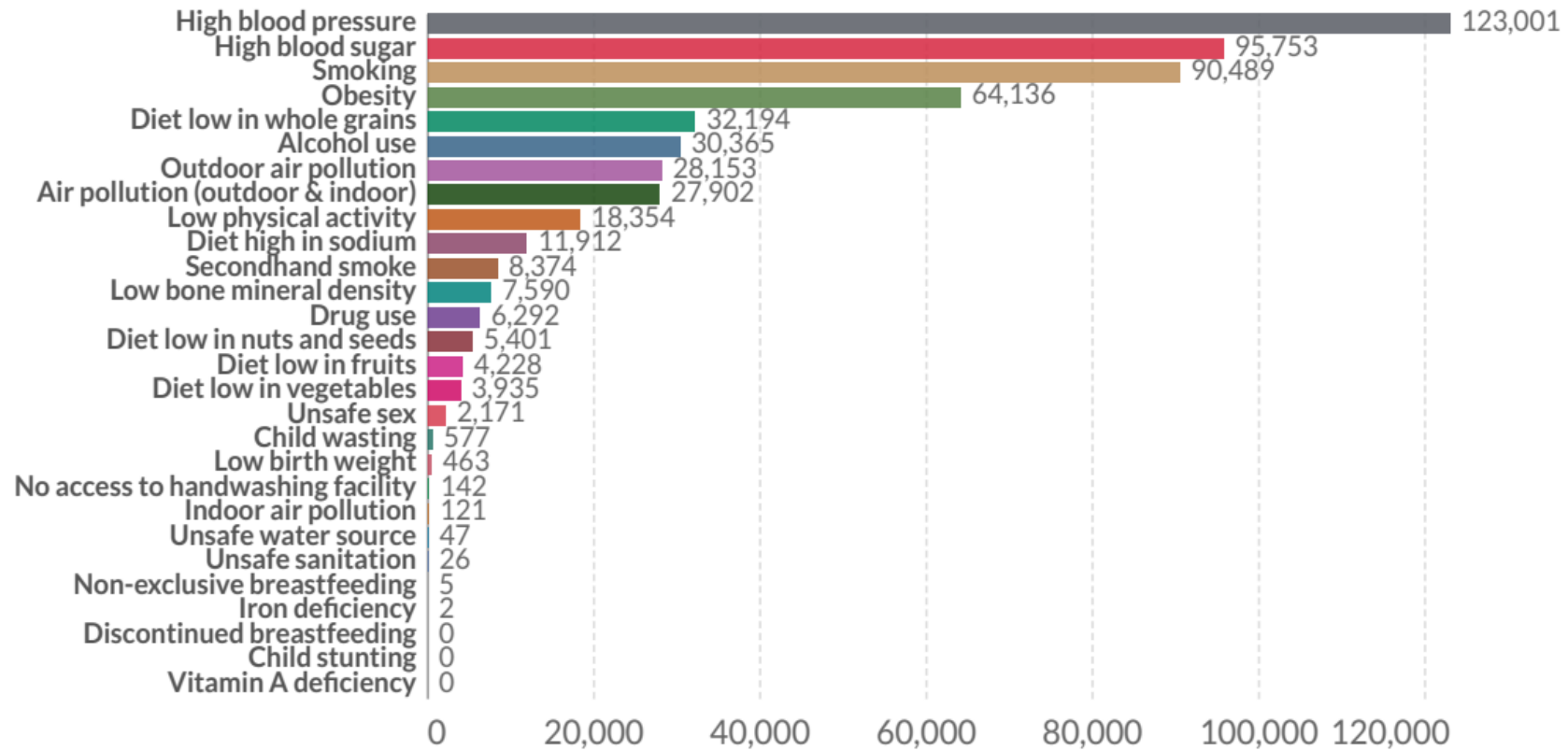
Figura 2. Numero di decessi per fattore di rischio, Italia, 2019

Number of deaths by risk factor, Italy, 2019

Our World
in Data

Total annual number of deaths by risk factor, measured across all age groups and both sexes.

↔ Change country



Source: IHME, Global Burden of Disease (GBD)

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Figura 3. Tasso di mortalità per inquinamento atmosferico, Mondo, 1990- 2019

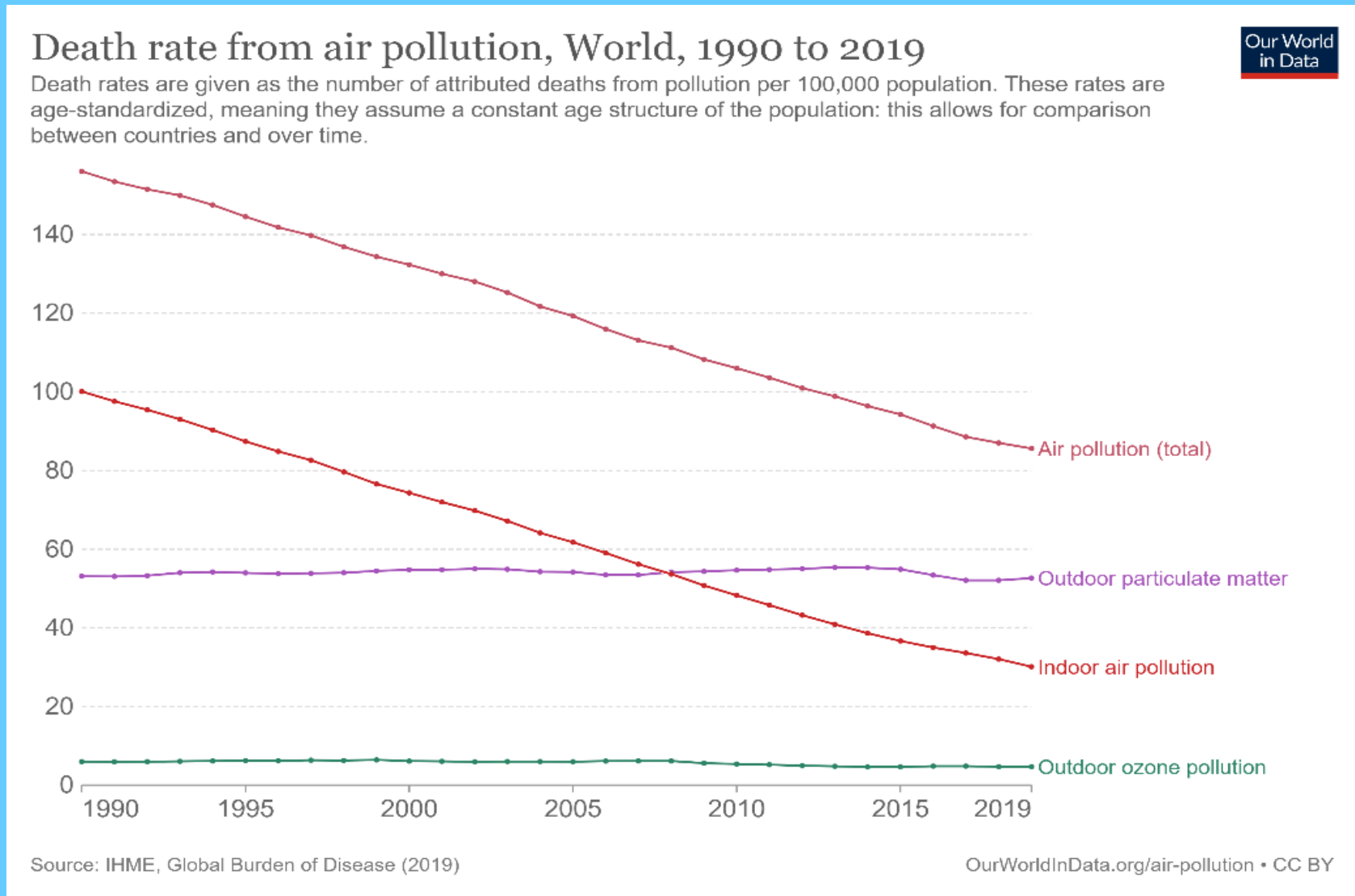
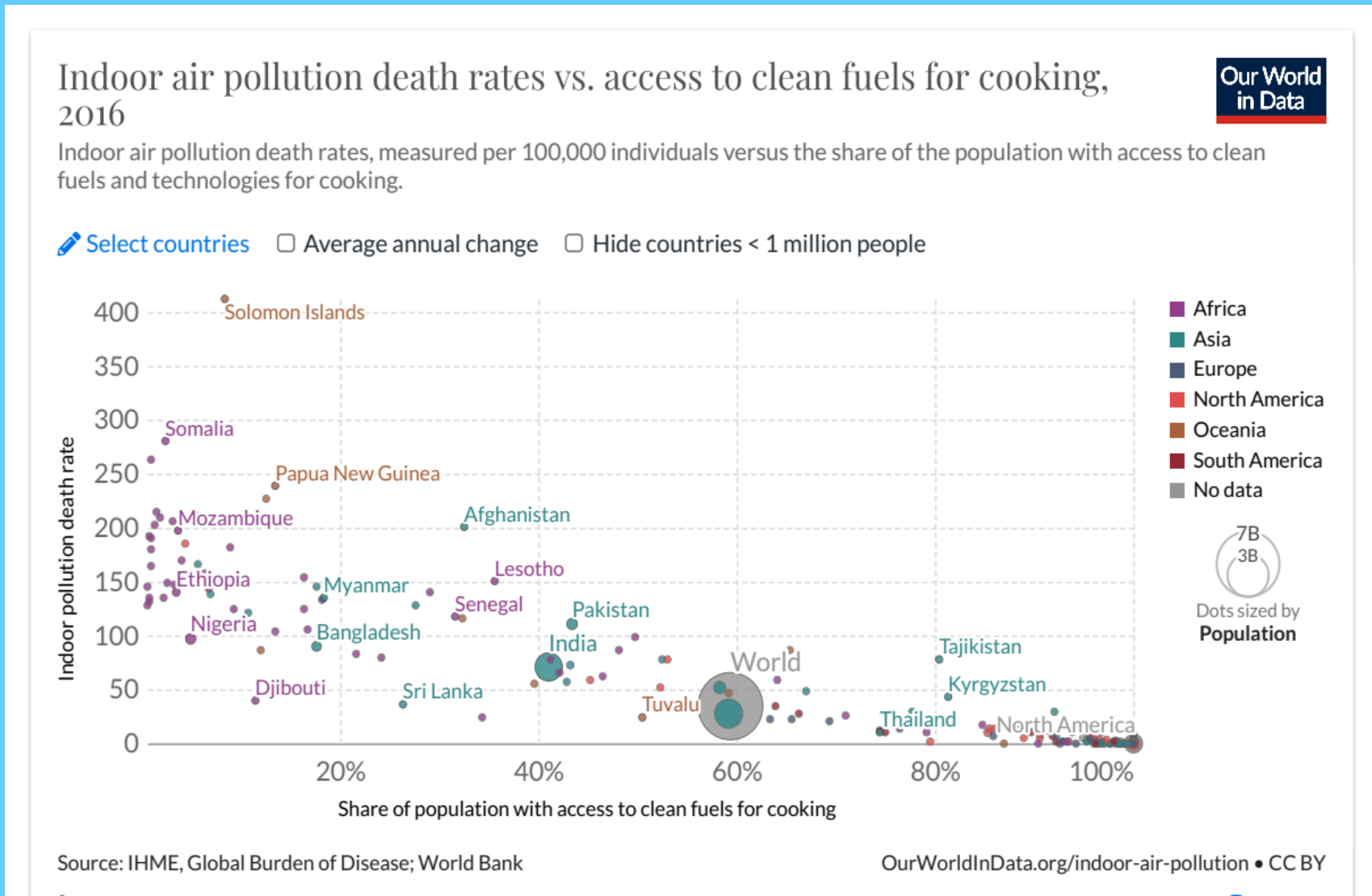


Tabella 6. Principali agenti indoor e potenziali fonti interne⁴⁵

FONTI	INQUINANTI
Processi di combustione a gas o carbone per riscaldare e/o cucinare, camini e stufe a legna, gas di scarico veicoli	Prodotti di combustione (CO, NO _x , SO ₂ , particolato)
Materiali da costruzione e isolanti	Amianto, fibre vetrose artificiali, particolato, Radon; Agenti biologici (per presenza di umidità e/o polvere)
Materiali di rivestimento e moquette	Formaldeide, acrilati, COV e Agenti biologici (per presenza di umidità e/o polvere)
Arredi	Formaldeide, COV e Agenti biologici (per presenza di umidità e/o polvere)
Liquidi e prodotti per la pulizia	Alcoli, fenoli, COV
Fotocopiatrici	Ozono (O ₃), polvere di toner, idrocarburi volatili (COV)
Fumo di sigaretta	Idrocarburi policiclici, COV, formaldeide, CO, particolato fine
Impianti di condizionamento	CO ₂ e COV (per scarso numero di ricambi orari o eccesso di riciclo); Agenti biologici (per mancanza di pulizia / manutenzione)
Polvere	Agenti biologici (allergeni indoor: acari)
Individui	CO ₂ e Agenti biologici (batteri, virus ecc.)
Animali	Allergeni indoor (peli, etc.)
Sorgenti naturali (lave, tufi, graniti, ecc.)	Radon

Figura 4. Tasso di mortalità per Inquinamento atmosferico indoor vs accesso a combustibile pulito per cucinare



Il raddoppio delle
concentrazioni
atmosferiche di CO₂(da
200 a700 (μL L⁻¹ induce
l'incremento della
produzione di polline
allergenico di ambrosia
(ragweed) del 61%

(Wayne P. Annals of Allergy,
Asthma & Immunolog, 2002)

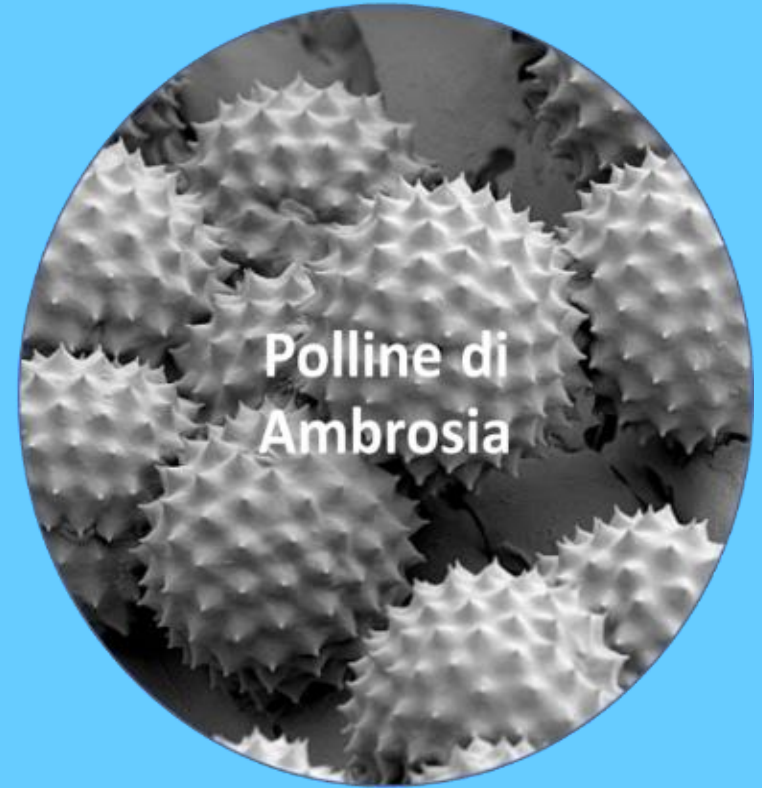


Tabella 7 *Interventi per la prevenzione delle malattie respiratorie croniche (MRC), con particolare riguardo al fattore di rischio indoor*


LIVELLO INDIVIDUALE	Aumentare la consapevolezza sull'utilità dell'uso continuo di inalatori per massimizzarne i benefici.
	Evitare l'uso della biomassa per cucinare o riscaldare e l'esposizione al fumo di seconda mano (ETS).
LIVELLO COMUNITARIO	Cambiare le percezioni sulle MRC e de-stigmatizzarle attraverso interventi educativi.
	Aumentare la consapevolezza sull'importanza della famiglia nel fornire supporto fisico, economico e psicologico ai pazienti con MRC.
LIVELLO DEL SISTEMA SANITARIO	Aumentare la comunicazione con i pazienti sul decorso della malattia e sui fattori di rischio.
	Migliorare le interazioni con la comunità attraverso operatori sanitari in prima linea.
	Coinvolgere la comunità con mezzi e metodi culturalmente accettabili e disponibili a livello locale.
	Migliorare la disponibilità e l'accessibilità della spirometria nelle cure primarie.
	Laddove la spirometria non è disponibile, prendere in considerazione alternative come questionari e strumenti validati (es. PEF), in particolare nei Paesi a basso-medio reddito.
	Attuare linee guida operative di assistenza primaria e programmi di formazione per operatori sanitari clinici e operatori sanitari di comunità.
	Migliorare la disponibilità e dell'accessibilità dei farmaci.
	Investire in strategie per ridurre gli ostacoli a comportamenti sani dei pazienti con CRD.
	Attuare programmi potenziati per combustibili più puliti e apparecchi migliori.
LIVELLO POLITICO	Sviluppare una politica sanitaria specifica per le MRC con protocolli basati sull'evidenza, supportati dalla regolare disponibilità e accessibilità di attrezzature mediche e farmaci.
	Definire norme specifiche per la gestione della qualità dell'aria interna.
LIVELLO DELLE SOCIETA' SCIENTIFICHE	Aumentare la consapevolezza sul carico socio-sanitario dei pazienti con CRD e sui relativi fattori di rischio.
	Lavorare insieme per perseguire la visione GARD di "un mondo in cui tutte le persone respirano liberamente".

Tabella 8 Valori di riferimento OMS


Inquinante	Tempo	Interim target				Livelli AQG
		1	2	3	4	
PM _{2,5} µg/m ³	Annuale	35	25	15	10	5
	24 h	75	50	37.5	25	15
PM ₁₀ µg/m ³	Annuale	70	50	30	20	15
	24 h	150	100	75	50	45
O ₃ µg/m ³	Picco stagionale	100	70			60
	8 h	160	120			100
NO ₂ µg/m ³	Annuale	40	30	20		10
	24 h	120	50			25
SO ₂ µg/m ³	24 h	125	50			40
CO mg/m ³	24 h	7				4

A series of narrative reviews on air pollution and respiratory health for Pulmonology: Why it is important and who should read it

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REVIEW
Issue 1 - “Update on adverse respiratory effects of outdoor air pollution”. Part 1): Outdoor air pollution and respiratory diseases: A general update and an Italian perspective
S. De Matteis^{a,b}, F. Forastiere^{c,d}, S. Baldacci^e, S. Maio^e, S. Tagliaferro^e, S. Fasola^d, G. Cilluffo^d, S. La Grutta^d, G. Viegi^{d,e,*}

Pulmonology. 2022 Jul-Aug;28(4):284-296

Abstract

Objective: to summarize the main updated evidence about the health effects of air pollution and to focus on Italian epidemiological experiences on the respiratory effects.

Results: the recent literature indicates that there is strong evidence for causal relationships between PM_{2.5} air pollution exposure and all-cause mortality as well as mortality from acute lower respiratory infections, ischaemic heart disease, stroke, chronic obstructive pulmonary disease, and lung cancer. A growing body of evidence also suggests causal relationships with type II diabetes and impacts on neonatal mortality from low birth weight and short gestation as well as neurologic effects in both children and adults. Italy, a Southern European country, faces a more threatening air pollution challenge because of the effects of both anthropogenic pollutants and natural dust (particulate matter, PM). The 2020 Report of the European Environment Agency highlighted the number of premature deaths in Italy attributable to main pollutants: 52,300 for PM_{2.5}, 10,400 for NO₂ and 3,000 for O₃ in 2018. In Italy, original time series and analytical epidemiological studies showed increased cardio-respiratory hospital admissions and mortality and increased risk of respiratory diseases in people living in urban areas.

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Outdoor air pollution and respiratory health

S. Maio, G. Sarno, S. Tagliaferro, F. Pirona, I. Stanisci, S. Baldacci, G. Viegi

The need to address the impact of air pollution on health is reinforced by recent scientific evidence and the 2021 WHO Air Quality Guidelines (AQG). Air pollution is an avoidable risk factor causing a high burden for society with elevated deaths, health disorders, disabilities and huge socio-economic costs, especially in low- and middle-income countries. We have evaluated recent evidence from international reports, systematic reviews and official websites of international agencies. Growing evidence shows a causal relationship between air pollution exposure and acute lower respiratory infections, chronic obstructive pulmonary disease, asthma and lung cancer. Exposure to air pollution in both the

short- and long-term has a serious impact on respiratory health. Harmful effects occur even at very low pollutant concentration levels, and there are no detectable thresholds below which exposure may be considered safe. The adverse respiratory health effects of air pollutants, even at low levels, are confirmed by recent epidemiological studies. Scientific respiratory societies and patient associations, along with other stakeholders in the health sector, should increase their engagement and advocacy to raise awareness of clean air policies and the latest WHO AQG.

KEY WORDS: particulate matter; PM_{2.5}; PM₁₀; asthma; COPD



“Grazie per l’invito e l’attenzione”.

Giovanni Viegi

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