

Original research

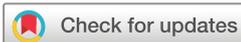
Occupational exposure to inorganic dust
and risk of gout: a population-based
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ABSTRACT

Background Risk factors operating independently of hyperuricemia could be of importance in determining why only a minority of people with hyperuricemia develop gout. Exposure to inorganic dust has been linked to other inflammatory diseases and could influence the development of gout.

Objectives To evaluate if occupational exposure to inorganic dust increases the risk of gout.

Methods Individuals aged 30–65 years with a first gout diagnosis in 2006–2012 in the population-based healthcare database of the Western Swedish Healthcare Region (VEGA) and population controls matched by age and sex were included. Data on occupation was collected from the Swedish occupational register. Exposure status was assigned by means of a job exposure matrix. Data on gout-related comorbidities was collected from VEGA. Alcohol use disorder and obesity were related both to gout and exposure to inorganic dust and were adjusted for in multivariate analyses. ORs were calculated using conditional logistic regression.

Results 5042 gout cases and 20 682 controls were included. Exposure to inorganic dust was associated with gout in both unadjusted (OR 1.12, 95% CI 1.04 to 1.20) and multivariate (OR 1.08, 95% CI 1.00 to 1.16) analyses of the whole population. In sex-stratified multivariate analyses, dust exposure was significantly associated with gout in women (adjusted OR 1.26, 95% CI 1.05 to 1.51), but not in men (adjusted OR 1.05, 95% CI 0.97 to 1.13).

Conclusions We describe for the first time an association between exposure to inorganic dust and gout. After adjusting for confounders, the findings were statistically significant for women but not for men.

INTRODUCTION

It is well established that hyperuricemia is a prerequisite of gout and that the risk of clinical gout increases in the presence of factors that increase serum urate levels, such as renal disease,^{1 2} alcohol use disorder,^{3 4} diuretic use⁵ and obesity.⁶ Some of these urate-elevating factors, such as alcohol use disorder and obesity, are associated with lower socioeconomic status (SES) as measured by income and level of education.⁷ This could in part explain the observation that gout is more common among people of lower SES. A number of epidemiological studies have found an increased prevalence of gout

Key message**What is already known about this subject?**

- Exposure to inorganic dust has been shown in several studies to associate with a number of rheumatic inflammatory diseases such as rheumatoid arthritis and systemic sclerosis, but it has not been investigated previously whether exposure is associated with an increased risk of gout.
- Inorganic dust, like silica, can activate the NLRP3 inflammasome, a key component in the inflammatory response to urate crystals, subsequently leading to clinical gout.

What does this study add?

- We demonstrate for the first time an association between occupational exposure to inorganic dust and incident gout.
- The findings are indicative of a previously unknown possible aetiological avenue for gout, in addition to known risk factors like genetics, diet and urate-elevating comorbidities.

How might this impact on clinical practice?

- If confirmed, the findings could provide a possibility for prevention of gout among susceptible individuals by minimising exposure to inorganic dust in the work environment.

among people with shorter education, lower income and manual occupations,^{8 11} where the difference is not fully explained by different rates of urate-elevating comorbidities or serum urate levels. The mechanism behind lower SES predisposing to gout independently of urate levels or urate-elevating comorbidities is unknown. One possible explanation is that yet unknown occupational exposures increasing the risk for gout could be more common in lower socioeconomic groups. Very little is known about the possible influence of occupational exposures on the risk of gout in recent times. It is possible that, yet unknown, exposures present in the work environment of today could increase the susceptibility to gout, by accenting either hyperuricemia, the process of

monosodium urate (MSU) crystallisation or the inflammatory response to MSU crystals.

Exposure to inorganic dusts, for example, silica dust, is one such occupational exposure that may be related to gout. Silica dust has been shown to activate the NLRP3 inflammasome,¹² which is a key component of the inflammatory response to MSU crystals.¹³ Exposure to silica has also been associated with renal disease,¹⁴ which could facilitate the development of hyperuricemia and gout. Exposure to inorganic dust has recently been shown in several studies to associate with a number of other rheumatic inflammatory diseases such as rheumatoid arthritis and systemic sclerosis.^{15–18} Increased levels of ambient air pollution, especially regarding the component of air pollution consisting of fine particulate matter <10 µM, have been reported to increase the occurrence of acute gout flares,¹⁹ but no studies investigating the possible association of occupational exposure to inorganic dust with gout are available.

The aim of the present study was therefore to examine if previous occupational exposure to inorganic dust predicted an incident diagnosis of gout, both overall and stratified by sex.

MATERIALS AND METHODS

Study design

This is a population-based nested case–control study including all individuals with a first diagnosis of gout in the years 2006–2012 and matched population controls without gout. Cases were matched to controls by age, sex and place of residence on municipality level. Up to five controls were matched to each individual with gout. In Sweden, each resident has a unique personal identification number that identifies records in official registers, enabling linking of data from different sources. For this study, we used data from several national and regional registers (described below) to identify individuals with gout and population controls and to define occupational exposure to inorganic dust as well as information on relevant confounders for the analyses.

Setting and study population

The Swedish healthcare system is tax-funded and offers universal access. Gout is typically diagnosed and treated in a primary care setting. All inhabitants in the Western Swedish Healthcare Region (WSHCR) from January 1, 2006 to December 31, 2012 constituted the study population. The population of WSHCR is approximately 1.6 million (20% of the total population of Sweden),²⁰ and the region is considered to be representative for Sweden with regard to health status, healthcare-seeking and socioeconomics.^{20–21}

Register sources

Western Swedish Healthcare Register (VEGA)

The register contains information about all healthcare contacts in primary care and inpatient and outpatient secondary care clinics in the WSHCR. Date of contact

and primary and secondary diagnoses given by the treating physician according to the Swedish version of the International Statistical Classification of Diseases (ICD) are registered. VEGA was used to identify gout cases and to retrieve information about relevant comorbidities for cases and controls. For gout cases, the date of the first gout diagnosis was used as the index date/date of identification.

Swedish Census Register

The census register holds demographic information about all registered inhabitants of Sweden including dates of death and emigration. The census register was used to identify up to five controls for each case, matched for age, sex and place of residence on municipality level at the date of identification. Controls were assigned the same index date/date of identification as their corresponding case.

Longitudinal Integration Database for Health Insurance and Labour Market Studies

The register is administered by Statistics Sweden. The database holds annual registers since 2001 on all individuals from 16 to 64 years of age, including information about occupation, employment status and educational level. Occupation is registered according to The Swedish Standard Classification of Occupations (SSYK) classification, which is based on the International Standard Classification of Occupations-88.²² Data on occupation is provided by employers. Complete data on occupation was available for between 91% and 94% of the population during the years 2001–2008.³⁵

Swedish Prescribed Drug Register

All prescribed drugs dispensed by Swedish pharmacies are recorded in the Swedish Prescribed Drug Register. The register has been available since July 1, 2005 and was used to determine dispensation of urate lowering therapy (ULT) and diuretics.

Case definition and selection of controls

To select individuals most likely to have new-onset gout, we included individuals with a diagnosis of gout in the VEGA register during 2006–2012. The VEGA database was then searched for gout diagnoses back to January 1, 2000, and individuals with a prior diagnosis of gout were excluded. All included individuals thus have a period free from gout diagnosis of at least 5 years before their index date. Gout diagnosis was defined by the presence of an ICD-10 code for gout (M10), registered at a visit to a physician in the VEGA database. We excluded cases and controls with dispensation of ULT- prior to first gout diagnosis or corresponding date for controls. The case definition has been previously validated by means of record review that showed a high validity of ICD-10 codes for gout in the VEGA database.²³

To ensure that the population included was part of the workforce, we included only individuals that were of

working age (30–65 years) at the time of gout diagnosis and that had at least one registered occupation during the 5 years prior to the year of identification with gout.

We applied the same conditions to the population controls, excluding controls with no occupation registered for the 5 years prior to the year of identification as control. The selection of cases and controls for the study is further described in online supplementary table 1.

Exposure

The occupational register was searched for registered occupational codes during the 5 calendar years prior to identification as case or control, that is, occupations on file in the occupational register for the years 2002–2006 contributed to exposure assignment for an individual that was identified as case or control in 2007. A job exposure matrix (JEM) for inorganic dust previously developed was used to assess exposure.²⁴ During development of the JEM, all occupations were re-assessed by two senior occupational hygienists as being occupationally exposed or not to inorganic dust including silica dust as previously described.^{24 25} An exposed occupation was defined based on the assumption that at least half of the subjects with these specific codes should have a strong probability of being exposed.

Individuals that had been employed in a dust-exposed occupation for at least 1 year during the 5-year-long predictor period were defined as exposed.

Definition of comorbidities

Comorbidities known to be associated with gout were identified in VEGA by ICD-10 codes. VEGA was searched from 2000 to 2012 for predefined comorbidities (for definitions used, see online supplementary table 2). A comorbidity was considered present if there was a healthcare contact with a corresponding ICD-10 coded diagnosis by a physician before the index date.

Statistics

The association between occupational exposure to inorganic dust and incident gout diagnosis was assessed using conditional logistic regression, accounting for the matched design of the study. Results are presented as unadjusted and adjusted ORs with 95% CIs, for the whole population and stratified by sex.

The covariates adjusted for in the analysis (alcohol use disorder and obesity) were chosen on the basis of being plausible confounders, that is, related to the outcome of gout based on prior clinical knowledge^{3 4 26} through the mechanism of elevating urate levels and associated with exposure to inorganic dust in our data (online supplementary tables 3–4). Other gout-related urate-elevating comorbidities, for example, renal disease and psoriasis, were not included in the multivariable analysis as they have no known association with occupational exposure to inorganic dust nor were they associated with dust exposure when this was tested in our own data (online supplementary tables 5–6).

In a secondary model, in addition to the covariates described above, we also adjusted for educational level.

RESULTS

Baseline characteristics of cases and controls

A 5042 gout cases and 20 682 population controls fulfilling the case definition were identified and included for analysis. Baseline characteristics of included individuals are shown in table 1.

Exposure to inorganic dust

Among individuals with incident gout, 30.1% were occupationally exposed to inorganic dust during the 5-year period preceding the index date, whereas 28.2% of population controls were exposed (p=0.006).

Among men, 34.4% of gout cases and 33.0% of the population controls were exposed to inorganic dust, corresponding to 1.4% (95% CI 0.2% to 3.1%) higher exposure frequency among the gout cases. In women, 16.0% of gout cases and 12.7% of controls were exposed, corresponding to 3.3% (95% CI 1.2% to 5.5%) higher exposure frequency among the gout cases (figure 1).

Association of exposure to inorganic dust to incident gout

Exposure to inorganic dust was associated with incident gout in the unadjusted analysis of the whole population (OR 1.12, 95% CI 1.04 to 1.20). In the multivariate analysis, after adjustment for alcohol use disorder and obesity, exposure to inorganic dust increased the odds of incident gout by 8% (OR 1.08, 95% CI 1.00 to 1.15) (figure 2A). In the results of the secondary model, where educational

Table 1 Baseline characteristics of included cases and controls

Variable	Gout cases, n=5042	Population controls, n=20 682
Men	3862 (76.6)	15 747 (76.1)
Age, mean(SD)	52.9 (8.9)	52.7 (8.9)
Education level >12 years	1283 (25.4)	6912 (33.4)
Exposed to inorganic dust during predictor period	1519 (30.1)	5822 (28.2)
Comorbidities		
Hypertension	2096 (41.6)	4474 (21.6)
Treatment with diuretics	1350 (26.8)	1969 (9.5)
Obesity	499 (9.9)	671 (3.2)
Diabetes	405 (8.0)	878 (4.2)
Renal disease	357 (7.1)	659 (3.2)
Coronary heart disease	263 (5.2)	484 (2.3)
Atrial fibrillation	242 (4.8)	314 (1.5)
Alcohol abuse	253 (5.0)	431 (2.1)
Thromboembolism	210 (4.2)	452 (2.2)
Congestive heart failure	196 (3.9)	138 (0.7)
Psoriasis	174 (3.5)	518 (2.5)
Stroke	93 (1.8)	235 (1.1)
Chronic obstructive pulmonary disease (COPD)	71 (1.4)	116 (0.6)

Except where otherwise indicated, values are no. (%).

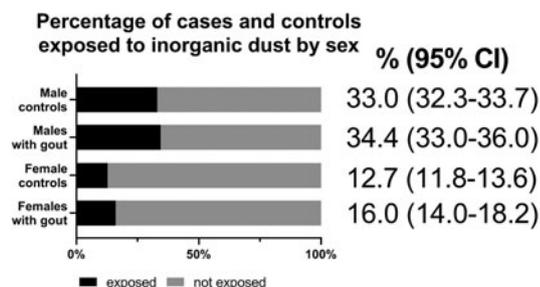


Figure 1 Percentage of male and female cases and controls exposed to inorganic dust during the 5 years before gout diagnosis or corresponding time period for controls.

level was also adjusted for, the association between inorganic dust and gout was diminished (**figure 2B**).

In the stratified analysis where men and women were analysed separately, exposure to inorganic dust

significantly increased the odds of gout for women by 32% in the unadjusted analysis (OR 1.32, 95% CI 1.10 to 1.58) and 26% in the adjusted analysis (OR 1.26, 95% CI 1.05 to 1.51). For men, the increase in odds of incident gout from exposure to inorganic dust was non-significant, OR 1.05, 95% CI 0.97 to 1.13 in the adjusted analysis (**figure 3**).

Most common occupations

In total, 7341 individuals (6526 men and 815 women) were defined by the JEM as exposed to inorganic dust. The five most common occupations among the exposed are shown in **table 2**.

The most common occupations defined as exposed differed for men and women; **table 3** shows the most common dust-exposed occupations for men and women separately.

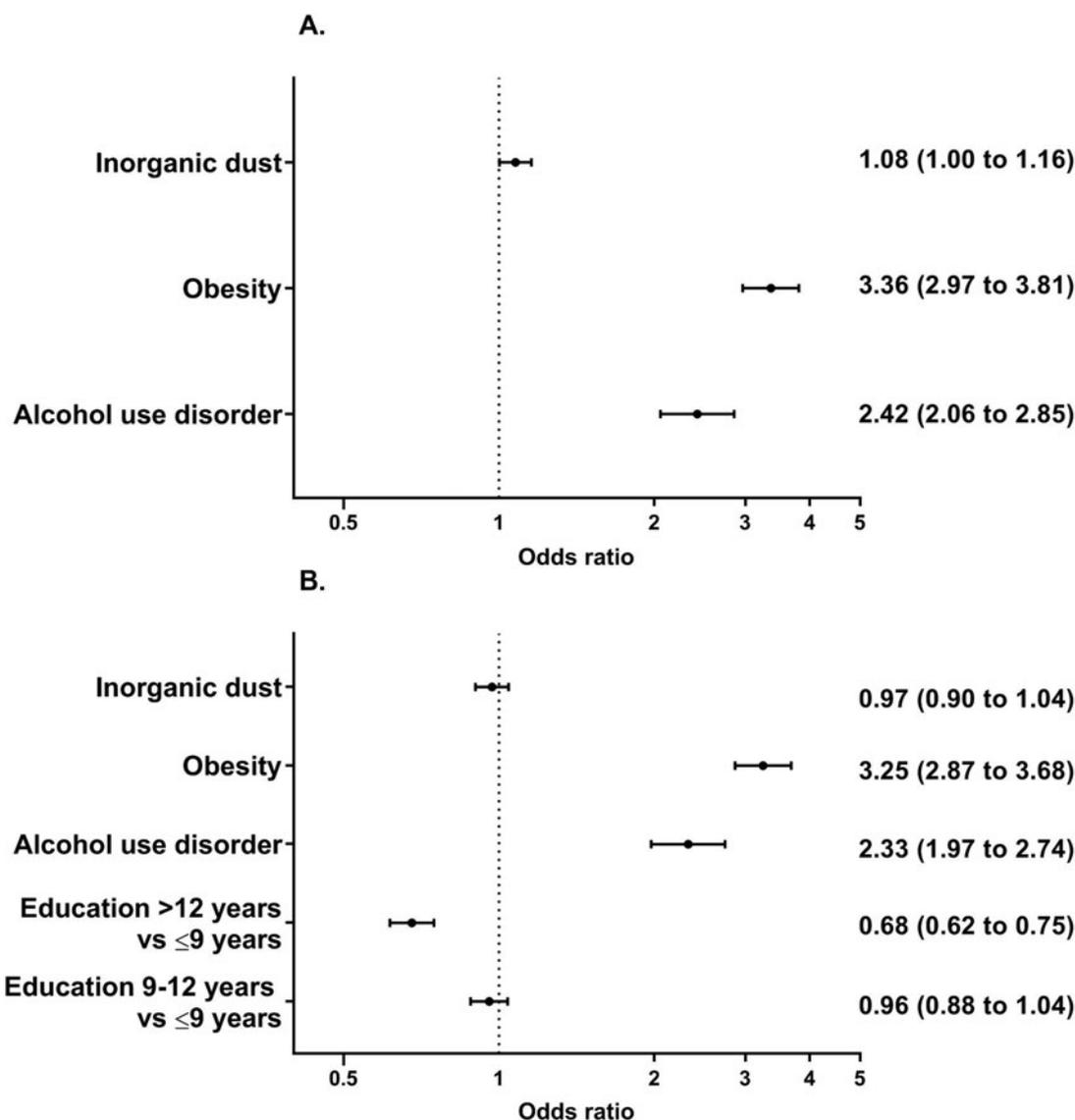


Figure 2 Predictors of incident gout. Multiadjusted odds ratios with 95% confidence intervals. A) Main model, adjusted for obesity and alcohol use disorder. B) Secondary model, adjusted for educational level in addition to obesity and alcohol use disorder.

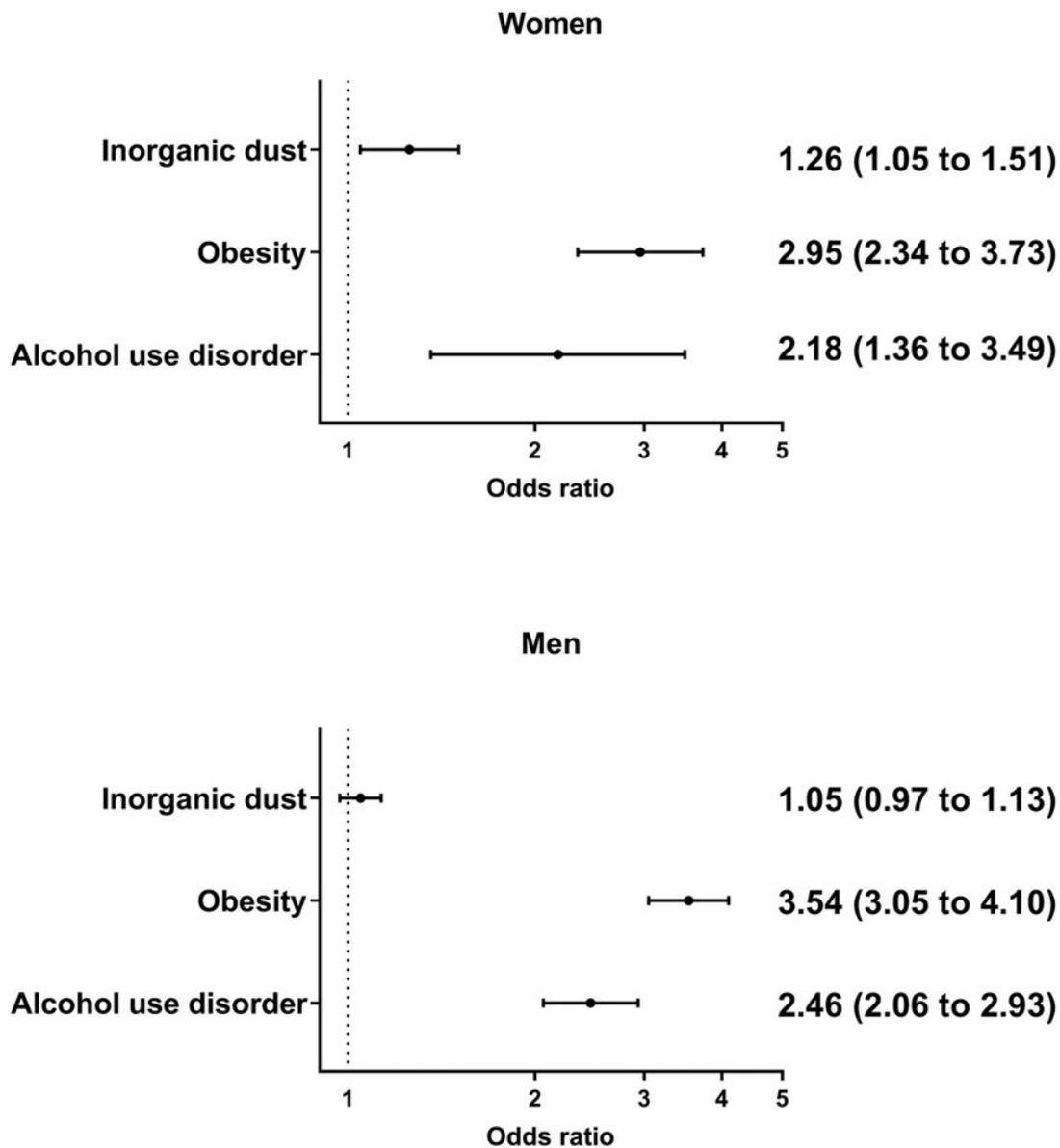


Figure 3 Predictors of incident gout in women and men. Multiadjusted odds ratios with 95% confidence intervals.

DISCUSSION

In the present study, we found that occupational exposure to inorganic dust was associated with incident gout. The association was statistically significant in women also after adjusting for confounding by diagnosed alcohol use

disorder and obesity. The effect was of modest size and smaller than what has been shown in studies of dust exposure and risk of rheumatoid arthritis and systemic sclerosis.^{15 16 18}

The pathogenetic mechanism through which inorganic dust exposure might increase the risk of gout is not clear. We speculate that a possible explanation could be the phenomenon of innate immune system training,²⁷ where priming of the innate immune system by prior inflammatory stimuli can cause increased reactivity later on. Occupational exposure to inorganic dust could possibly trigger innate immune system training through activation of the NLRP3 inflammasome,¹² leading to increased interleukin-1 production and in the long run to increased responses of the innate immune system towards urate crystals, manifesting as increased incidence of clinical gout in dust-exposed individuals.

Table 2 Most common occupations exposed to inorganic dust

Occupational title	N (%)*
Building finishers and related trades workers	1081 (15)
Motor vehicle drivers	672 (9)
Material recording and transport clerks	633 (9)
Assemblers	549 (7)
Building frame and related trades workers	486 (7)

*Percentages refer to % of all exposed individuals.

Table 3 Most common occupations defined as exposed for men and women separately

Women—5 most common occupations defined as exposed		Men—5 most common occupations defined as exposed	
Job title	n (%)*	Job title	n (%)†
Domestic and related helpers, cleaners and launderers	455 (56)	Building finishers and related trades workers	1060 (16)
Building caretakers, window and related cleaners	58 (7)	Motor vehicle drivers	669 (10)
Assemblers	54 (7)	Material recording and transport clerks	587 (9)
Material recording and transport clerks	46 (6)	Assemblers	495 (8)
Other machine operators not elsewhere classified	43 (5)	Building frame and related trades workers	483 (7)

*Percentages refer to percent of all exposed women.

†Percentages refer to percent of all exposed men.

Studies on associations between air pollution and gout occurrence are scarce, but our findings of an association between dust exposure and gout are in line with a previous report from Korea, where increased levels of air pollution with fine particulate matter were associated with acute gout flares.¹⁹

Hyperuricemia and factors, like obesity and alcohol use disorder, which contribute to hyperuricemia and clinical gout are by far the most important risk factors for gout. In relation to such factors, the effects of occupational exposures are small but, as we have shown, possibly important. Uncovering novel risk factors for gout unrelated to urate metabolism could contribute to an increased understanding of the pathogenesis and help explain why only a minority of individuals with hyperuricemia develop clinical gout and in the long run possibly contribute to prevention of gout in susceptible individuals.

Other occupational exposures could also play a role in the development of clinical gout and should be investigated in further studies. There is considerable evidence that lead exposure, even when not leading to overtly elevated blood levels of lead, influences urate levels.^{28 32} Consequently, it should be investigated in future studies whether occupations that entail exposure to lead increase the risk of clinical gout.

There are some limitations of the study that should be discussed and acknowledged. First, there could be misclassification of exposure based on occupational coding. The validity of the occupational codes reported to the occupational register by employers has not been investigated specifically for dust-exposed occupations. In our data, occupational codes on the individual level were very consistent over the time period of 5 years, strongly

suggesting that the quality of reporting of occupational codes is good. Furthermore, any occupational misclassification is likely to be non-directional, as misclassification should be equally common for cases and controls.

Second, as in all observational register studies, residual confounding could be present.

Third, misclassification of comorbidities could be a problem as they were defined by the presence of ICD-10 codes. In general, the validity of ICD-10 codes in the Swedish National Patient Register is good.³³ However, all diagnoses have not been validated, and under-reporting of, for example, alcohol use disorder and obesity is likely to be present, which could influence our results. Such under-reporting is however likely to be of equal magnitude for both cases and controls.

Finally, adjustment for additional possible confounders should be discussed. Whether to adjust for educational level is a complicated issue when assessing potential effects of occupational exposures. On the one hand, educational level is a marker of SES, which has an influence on the health of individuals. On the other hand, educational level is strongly connected to choice of occupation. Choice of occupation can thus be seen to be in the temporal pathway between education and clinically evident gout. Low educational level strongly predicts having a dust-exposed job. It seems unlikely that educational level in itself should cause disease. Rather, downstream effects of a short education, such as ending up in a dust-exposed job, could have biological effects and increase the likelihood of disease. Thus, adjustment for educational status could be considered to be overadjustment, that is, adjusting for an upstream mediator of the variable of interest. In line with this, in the model where educational level was adjusted for, the effect of inorganic dust exposure on risk of gout was no longer evident. This finding could either be an effect of overadjustment, or possibly it shows that socioeconomic factors, of which educational level is a marker, are of greater importance than occupational exposure to inorganic dust for the risk of gout.

Furthermore, smoking exposure could not be included in the present analyses, but although it is a risk factor for many diseases, it has not consistently been shown to associate with urate levels or clinical gout.³⁴ Nonetheless, residual confounding due to smoking cannot be excluded.

There are also several strengths of the study. First, the population-based design, including all individuals with an incident gout diagnosis in the region, minimises the risk of selection bias. The validity of the case definition for gout used in the study has been studied and was found to be high.²³ Second, the use of coded occupational titles from an official register and a JEM to determine exposure is another strength of the study, as this avoids the potential recall bias associated with retrospective self-reported occupation.

In conclusion, we describe for the first time an association between occupational exposure to inorganic dust and incident gout.

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Competing interests None declared.

Patient consent for publication Not required.

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Provenance and peer review Not commissioned; externally peer reviewed.

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REFERENCES

- Roughley MJ, Belcher J, Mallen CD, *et al*. Gout and risk of chronic kidney disease and nephrolithiasis: meta-analysis of observational studies. *Arthritis Res Ther* 2015;17.
- Juraschek SP, Kovell LC, Miller ER, *et al*. Association of kidney disease with prevalent gout in the United States in 1988–1994 and 2007–2010. *Semin Arthritis Rheum* 2013;42:551–61.
- Tu H-P, Tung Y-C, Tsai W-C, *et al*. Alcohol-related diseases and alcohol dependence syndrome is associated with increased gout risk: a nationwide population-based cohort study. *Joint Bone Spine* 2017;84:189–96.
- Choi HK, Atkinson K, Karlson EW, *et al*. Alcohol intake and risk of incident gout in men: a prospective study. *Lancet* 2004;363:1277–81.
- Bruderer S, Bodmer M, Jick SS, *et al*. Use of diuretics and risk of incident gout: a population-based case-control study. *Arthritis Rheumatol* 2014;66:185–96.
- Rassen JA, Bartels DB, Schneeweiss S, *et al*. Measuring prevalence and incidence of chronic conditions in claims and electronic health record databases. *Clin Epidemiol* 2018;11:1–15.
- Public Health Agency of Sweden. Öppna Jämförelser Folkhälsa 2019. 2019. Available from: <https://www.folkhalsomyndigheten.se/contentassets/ec714fca0b0145eab3d7924511550a74/oppna-jamforelser-folkhalsa-2019-18076.pdf> (accessed 13 02 2019)
- Gardner MJ, Power C, Barker DJ, *et al*. The prevalence of gout in three English towns. *Int J Epidemiol* 1982;11:71–5.
- Kapetanovic MC, Hameed M, Turkiewicz A, *et al*. Prevalence and incidence of gout in Southern Sweden from the socioeconomic perspective. *RMD Open* 2016;2:e000326.
- Hayward RA, Rathod T, Roddy E, *et al*. The association of gout with socioeconomic status in primary care: a cross-sectional observational study. *Rheumatology (Oxford)* 2013;52:2004–8.
- Vazquez-Mellado J, Cruz J, Guzman S, *et al*. Severe tophaceous gout. Characterization of low socioeconomic level patients from México. *Clin Exp Rheumatol* 2006;24:233–8.
- Peeters PM, Perkins TN, Wouters EF, *et al*. Silica induces NLRP3 inflammasome activation in human lung epithelial cells. *Part Fibre Toxicol* 2013;10:3.
- Martinon F, Petrioli V, Mayor A, *et al*. Gout-associated uric acid crystals activate the NALP3 inflammasome. *Nature* 2006;440:237–41.
- Maciejewska A. Health effects of occupational exposure to crystalline silica in the light of current research results. *Med Pr* 2014;65:799–818.
- Vihlborg P, Bryngelsson IL, Andersson L, *et al*. Risk of sarcoidosis and seropositive rheumatoid arthritis from occupational silica exposure in Swedish iron foundries: a retrospective cohort study. *BMJ Open* 2017;7:e016839.
- Ilar A, Alfredsson L, Wiebert P, *et al*. Occupation and risk of developing rheumatoid arthritis: results from a population-based case-control study. *Arthritis Care Res (Hoboken)* 2018;70:499–509.
- Too CL, Muhamad NA, Ilar A, *et al*. Occupational exposure to textile dust increases the risk of rheumatoid arthritis: results from a Malaysian population-based case: control study. *Ann Rheum Dis* 2016;75:997–1002.
- Blanc PD, Jarvholm B, Toren K. Prospective risk of rheumatologic disease associated with occupational exposure in a cohort of male construction workers. *Am J Med* 2015;128:1094–101.
- Ryu H, Seo M, Choi H, *et al*. THU0427 Ambient air pollution and risk of acute gout flares: a time-series study. *Ann Rheum Dis* 2017;76:369–
- Statistics Sweden. Folkmängden efter region, civilstånd, ålder och kön. År 1968–2014. 2012. Available from: https://www.statistikdatabasen.scb.se/pxweb/sv/ssd/START_BE_BE0101A/BefolkningNy/?rxid=35ac0e2c-0f41-4674-8730-34d07c219c88. (accessed 2020 06 29).
- Ministry of Health and Social Affairs SGO. Updated high-cost protection: outpatient care and medication.
- International Labour Office. *International Standard Classification of Occupations: ISCO-88*. Geneva: International Labour Office, 1991.
- Dehlin M, Stasinopoulou K, Jacobsson L. Validity of gout diagnosis in Swedish primary and secondary care – a validation study. *BMC Musculoskelet Disord* 2015;16:149.
- Lillienberg L, Andersson E, Janson C, *et al*. Occupational exposure and new-onset asthma in a population-based study in Northern Europe (RHINE). *Ann Occup Hyg* 2013;57:482–92.
- Toren K, Blanc PD, Naidoo RN, *et al*. Occupational exposure to dust and to fumes, work as a welder and invasive pneumococcal disease risk. *Occup Environ Med* 2020;77:57–63.
- Choi HK, Atkinson K, Karlson EW, *et al*. Obesity, weight change, hypertension, diuretic use, and risk of gout in men: the health professionals follow-up study. *Arch Intern Med* 2005;165:742–8.
- Moorlag S, Roring RJ, Joosten LAB, *et al*. The role of the interleukin-1 family in trained immunity. *Immunol Rev* 2018;281:28–39.
- Baki AE. The effects of lead exposure on serum uric acid and hyperuricemia in young adult workers: a cross-sectional controlled study. *Archives Rheumatol* 2016;31:71–5.
- Dai H, Huang Z, Deng Q, *et al*. The effects of lead exposure on serum uric acid and hyperuricemia in Chinese adults: a cross-sectional study. *Int J Environ Res Public Health* 2015;12:9672–82.
- Lee D, Choi W-J, Oh J-S, *et al*. The relevance of hyperuricemia and metabolic syndrome and the effect of blood lead level on uric acid concentration in steelmaking workers. *Annals Occup Env Med* 2013;25:27.
- Krishnan E, Lingala B, Bhalla V. Low-level lead exposure and the prevalence of gout: an observational study. *Ann Intern Med* 2012;157:233–41.
- Alasia DD, Emem-Chioma PC, Wokoma FS. Association of lead exposure, serum uric acid and parameters of renal function in Nigerian lead-exposed workers. *Int J Occup Environ Med* 2010;1:182–90.
- Ludvigsson JF, Andersson E, Ekbohm A, *et al*. External review and validation of the Swedish national inpatient register. *BMC Public Health* 2011;11:450.
- Fanning N, Merriman TR, Dalbeth N, *et al*. An association of smoking with serum urate and gout: a health paradox. *Semin Arthritis Rheum* 2018;47:825–42.
- Statistics Sweden. Background Facts. Labour and education statistics. 2011. Available from: https://www.scb.se/contentassets/f1ebee6d63544edb9a83148b5674271/am0208_2011a01_br_am76br1105.pdf (accessed: 2020 06 30).