

Perfluoroalkyl chemicals and elevated serum uric acid in US adults

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Background: Perfluoroalkyl chemicals, including perfluorooctanoic acid and perfluorooctane sulfonate, are man-made chemicals that have been detected in the blood of over 98% of the US population. Serum uric acid is a novel biomarker, even mild elevations of which has been implicated in the development of hypertension, diabetes mellitus, cardiovascular disease, and chronic kidney disease. We examined the relationship of serum perfluoroalkyl chemicals, including perfluorooctanoic acid and perfluorooctane sulfonate, and elevated uric acid levels in a representative sample of US adults.

Methods: We examined 3883 participants from the 1999–2000 and 2003–2006 National Health and Nutritional Examination Surveys, a representative, multiethnic population-based survey of noninstitutionalized US adults. Serum perfluorooctanoic acid and perfluorooctane sulfonate were analyzed as quartiles. The main outcome was hyperuricemia.

Results: We found that serum levels of perfluoroalkyl chemicals, including perfluorooctanoic acid and perfluorooctane sulfonate, were positively associated with hyperuricemia. This association appeared to be independent of confounders such as age, gender, race-ethnicity, body mass index, diabetes, hypertension, and serum cholesterol. Compared with subjects in quartile 1 (referent), the multivariate odds ratio for hyperuricemia among subjects in quartile 4 was 1.97 (95% confidence interval 1.44–2.70, $P < 0.0001$) for perfluorooctanoic acid and 1.48% (95% confidence interval 0.99–2.22, $P = 0.0433$) for perfluorooctane sulfonate. This observed association persisted in subgroup analysis by gender and body mass index.

Conclusion: Our results demonstrate that elevated levels of perfluoroalkyl chemicals are associated with hyperuricemia even at low perfluoroalkyl chemical exposure levels as seen in the US general population.

Keywords: perfluoroalkyl chemicals, perfluorooctanoic acid, perfluorooctane sulfonate, uric acid

Introduction

There is concern regarding human exposure to perfluoroalkyl chemicals, including perfluorooctanoic acid and perfluorooctane sulfonate, because these chemicals are persistent in the environment, bioaccumulated, biomagnified along food chains, and have been shown to cause developmental and other adverse health effects in laboratory animals.^{1,2} Perfluoroalkyl chemicals have been widely used in the manufacture of industrial and consumer products such as surfactants, lubricants, polishes, paper and textile coatings, food packaging, and fire-retarding foams.² Perfluoroalkyl chemicals have been detected in the blood of more than 98% of the US population.³

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Uric acid is a byproduct of purine metabolism which has both oxidant as well as antioxidant properties. Several studies have shown that higher serum uric acid levels are associated with markers of inflammation,⁴ insulin resistance,⁵ and elevated lipid levels.⁶ Studies have also reported an association between higher serum uric acid levels and risk of developing hypertension,⁷ diabetes mellitus,⁸ chronic kidney disease,⁹ and cardiovascular disease.¹⁰ High levels of serum uric acid is also considered to be the underlying metabolic derailment in gout.¹¹

Sakr et al^{12,13} and Costa et al¹⁴ in two separate cohorts of occupationally exposed workers reported that perfluorooctanoic acid levels are cross-sectionally associated with elevated uric acid. Recently, Steenland et al¹⁵ reported modest positive associations between both serum perfluorooctanoic acid and perfluorooctane sulfonate levels and elevated uric acid in the C8 Health Study, a community-based study of residents in six water districts in Ohio and West Virginia who were exposed to very high levels of perfluorooctanoic acid following contamination from a chemical plant. However, perfluorooctanoic acid exposure levels in the US general population are much lower than those reported in these studies which examined high-exposure populations. In this context, a recent report based on a community-based sample of subjects exposed to perfluorooctanoic acid emissions from an industrial facility reported a concentration-dependent clearance for perfluorooctanoic acid, with half-lives of 2.9 years at higher exposure levels and 8.5 years for lower exposure levels.¹⁶ Therefore, it is also important to study perfluorooctanoic acid exposure at lower serum levels because at such concentrations these chemicals may potentially persist in the body for longer. Therefore, we examined if there was a positive association between serum perfluoroalkyl chemicals and uric acid in the National Health and Nutritional Examination Survey (NHANES), a representative, multiethnic sample of the US general population.

Materials and methods

The current study is based on six years of combined data from the 1999–2000, 2003–2004, and 2005–2006 NHANES. Detailed description of the NHANES study design and methods are available elsewhere.¹⁷ In brief, the NHANES survey included a stratified, multistage probability sample representative of the civilian noninstitutionalized US population. Selection was based on counties, blocks, households, and individuals within households, and included the

oversampling of low-income persons, persons 60+ years, African Americans, and Mexican Americans in order to provide stable estimates of these groups. The survey also includes biomonitoring for different environmental chemicals, including perfluoroalkyl chemicals, in a random one third subsample of participants by the National Center for Environmental Health. Subjects were required to sign a consent form before their participation, and approval was obtained from the Human Subjects Committee at the US Department of Health and Human Service.

The current study sample consisted of 3974 participants aged ≥ 20 years who had perfluoroalkyl chemical measurements available. We excluded subjects with missing data ($n = 91$) on uric acid and covariates included in the multivariable model, including education level, body mass index, or cholesterol levels. This resulted in 3883 participants (51.7% women).

Exposure measurements

Age, gender, race/ethnicity, smoking status, alcohol intake (g/day), level of education, history of diabetes, oral hypoglycemic intake, insulin administration, and antihypertensive medication use were assessed using a questionnaire. Individuals who had not smoked ≥ 100 cigarettes in their lifetimes were considered never smokers; those who had smoked ≥ 100 cigarettes in their lifetimes were considered former smokers if they answered negatively to the question “Do you smoke now?” and current smokers if they answered affirmatively. Body mass index was calculated as weight in kilograms divided by height in meters squared. Heavy drinking was defined as consumption of more than two drinks/day for the question on an average alcohol intake over the past 12 months.

Rigorous procedures with quality control checks were used in blood collection, and details about these procedures are provided in the NHANES Laboratory/Medical Technologists Procedures Manual.¹⁸ Perfluoroalkyl chemicals were measured in serum by the National Center for Environmental Health using automated solid-phase extraction coupled to isotope dilution high-performance liquid chromatography-tandem mass spectrometry; details of laboratory methods are available elsewhere.³ Our study examined the perfluoroalkyl chemicals, perfluorooctanoic acid and perfluorooctane sulfonate, that were detected in greater than 98% of people. Values below the limit of detection were reported by NHANES as the limit of detection divided by the square root of 2.

Serum total cholesterol was measured enzymatically. Seated systolic and diastolic blood pressures were measured using a mercury sphygmomanometer according to the American Heart Association and Seventh Joint National Committee (JNC7) recommendations.¹⁹ Up to three measurements were averaged for systolic and diastolic pressures. Patients were considered hypertensive if they reported current blood pressure-reducing medication use and/or had systolic blood pressures ≥ 140 mmHg and/or diastolic blood pressures ≥ 90 mmHg.

Outcome of interest

The main outcome of interest was serum uric acid level or the presence of hyperuricemia. Uric acid was measured using the Beckman Synchron LX20 system. Details of laboratory measurement are available online.²⁰ In brief, uric acid is oxidized by uricase to produce allantoin and hydrogen peroxide. The hydrogen peroxide reacts with 4-aminoantipyrine and 3,5-dichloro-2-hydroxybenzene sulfonate in a reaction catalyzed by peroxidase to produce a colored product. The Beckman Synchron LX20 system monitors the change in absorbance at 520 nm at a fixed time interval. The change in absorbance is directly proportional to the concentration of uric acid in the sample. Coulston Foundation at Alamogordo, New Mexico, performed testing in 1999–2000 and Collaborative Laboratory Services at Ottumwa, Iowa, performed testing in 2003–2004. Hyperuricemia was defined as serum uric acid levels >6.8 mg/dL in men and >6.0 mg/dL in women.¹¹

Statistical analysis

Serum perfluoroalkyl chemicals, including perfluorooctanoic acid and perfluorooctane sulfonate, were analyzed both as a continuous as well as a categorical variable. For analysis as a continuous variable, perfluoroalkyl chemical values were log-transformed (base 2) as a result of their skewed distribution. We also categorized serum perfluoroalkyl chemicals into quartiles. We ran linear regression models with serum perfluoroalkyl chemical quartiles as the independent variable to examine the mean change in serum uric acid with increasing categories of perfluoroalkyl chemical, taking the lowest perfluoroalkyl chemical quartile as the referent. We ran three nested models, ie, the unadjusted model, the age and gender-adjusted model, and the multivariable-adjusted model, additionally adjusting for race-ethnicity (non-Hispanic whites, non-Hispanic blacks, Mexican Americans, others), education categories (<high school, high school, >high school), smoking (never, former, current), alcohol intake

(heavy drinker), body mass index (kg/m²), hypertension (absent, present), diabetes (absent, present), and serum total cholesterol (mg/dL). Subsequently, we ran multivariable logistic regression models to calculate the odds ratio ([OR], 95% confidence interval [CI]) of hyperuricemia for each higher perfluoroalkyl chemical level by taking the lowest category as the referent. Trends in the OR of hyperuricemia across increasing serum perfluoroalkyl chemical levels were determined by modeling perfluoroalkyl chemical categories as an ordinal variable. As recommended by the National Center for Health Statistics,²¹ sample weights that account for the unequal probabilities of selection, oversampling, and nonresponse in the NHANES survey were applied for all analyses. Analyses were conducted using SUDAAN (version 8.0, Research Triangle Institute, Research Triangle Park, NC) and SAS (version 9.2, SAS Institute, Cary, NC) software. Standard errors were estimated using the Taylor series linearization method.

Results

Table 1 presents the characteristics of the population. Overall, this was a middle-aged representative sample of the US population. It comprised 51.7% women,

Table 1 Characteristics of study population

Characteristics	Percentages or mean values \pm SE
Unweighted sample size	3883
Age, years	46.4 \pm 0.5
Women, %	51.7
Race-ethnicity, %	
Non-Hispanic whites	73.5
Non-Hispanic blacks	10.7
Mexican Americans	7.8
Others	8.0
Education categories, %	
Below high school	18.5
High school	27.1
Above high school	54.4
Smoking, %	
Never smoker	49.9
Former smoker	26.0
Current smoker	24.1
Alcohol intake, %	
Heavy drinker	24.3
Body mass index, kg/m ²	28.5 \pm 0.2
Total cholesterol, mg/dL	200.7 \pm 0.9
Diabetes mellitus, %	10.2
Hypertension, %	31.2
Serum uric acid, mg/dL	5.4 \pm 0.04
Hyperuricemia, %	19.2

Abbreviation: SE, standard error of the mean.

Table 2 Association between serum PFOA level and uric acid levels

Plasma PFC level*	Sample size	Unadjusted mean change in uric acid, mg/dL (95% CI)	Age, gender-adjusted mean change in uric acid, mg/dL (95% CI)	Multivariate-adjusted mean change in uric acid, mg/dL (95% CI)†
PFOA*				
Quartile 1	984	1 (referent)	1 (referent)	1 (referent)
Quartile 2	931	0.20 (0.05, 0.34)	0.15 (0.03, 0.27)	0.14 (0.04, 0.25)
Quartile 3	1003	0.40 (0.23, 0.56)	0.37 (0.23, 0.51)	0.37 (0.25, 0.49)
Quartile 4	965	0.56 (0.43, 0.68)	0.48 (0.35, 0.60)	0.44 (0.32, 0.56)
P trend		<0.0001	<0.0001	<0.0001
Log-transformed PFOA		0.45 (0.37, 0.53)	0.25 (0.17, 0.32)	0.22 (0.15, 0.30)
PFOS†				
Quartile 1	967	1 (referent)	1 (referent)	1 (referent)
Quartile 2	969	0.12 (-0.07, 0.30)	0.16 (-0.01, 0.32)	0.18 (0.05, 0.31)
Quartile 3	978	0.28 (0.10, 0.45)	0.24 (0.05, 0.43)	0.22 (0.04, 0.40)
Quartile 4	969	0.29 (0.11, 0.46)	0.26 (0.09, 0.44)	0.27 (0.13, 0.41)
P trend		0.0006	0.0051	0.0018
Log-transformed PFOS		0.40 (0.31, 0.50)	0.16 (0.05, 0.26)	0.14 (0.05, 0.23)

Notes: *Plasma PFOA quartiles: quartile 1 (<2.4 ppb), quartile 2 (2.4–3.4 ppb), quartile 3 (3.5–5.1 ppb), quartile 4 (>5.1 ppb); in men: quartile 1 (<3.2 ppb), quartile 2 (3.2–4.5 ppb), quartile 3 (4.6–6.4 ppb), quartile 4 (>6.4 ppb); †plasma PFOS quartiles: quartile 1 (<11.2 ppb), quartile 2 (11.2–17.8 ppb), quartile 3 (17.9–27.9 ppb), quartile 4 (>27.9 ppb); in men: quartile 1 (<17.5 ppb), quartile 2 (17.5–24.7 ppb), quartile 3 (24.8–35.6 ppb), quartile 4 (>35.6 ppb); ‡adjusted for age (years), gender (men, women), race-ethnicity (non-Hispanic whites, non-Hispanic blacks, Mexican Americans, others), education categories (<high school, high school, >high school), smoking (never, former, current), alcohol intake (heavy drinker), body mass index (kg/m²), hypertension (absent, present), diabetes (absent, present), and serum total cholesterol (mg/dL).

Abbreviations: CI, confidence interval; PFOA, perfluorooctanoic acid; PFOS, perfluorooctane sulfonate.

73.5% non-Hispanic whites, 10.7% non-Hispanic blacks, and 7.8% Mexican Americans. The mean serum uric acid level was 5.4 mg/dL, and 19.2% of the population had hyperuricemia.

Table 2 presents the association between increasing serum perfluorooctanoic acid, perfluorooctane sulfonate, and uric acid in the whole study sample. We observed a positive association between increasing quartiles of serum perfluorooctanoic acid and perfluorooctane sulfonate and a change in uric acid levels in the unadjusted, age-adjusted, and gender-adjusted models, as well as the multivariable-adjusted models. Models evaluating trend in this association were also statistically significant.

Tables 3 and 4 present the association between increasing serum perfluorooctanoic acid and perfluorooctane sulfonate and serum uric acid stratified by gender and body mass index categories, respectively. Similar to the results in Table 2, we observed a positive association between increasing quartiles of serum perfluorooctanoic acid and perfluorooctane sulfonate and change in uric acid levels. Models evaluating trend in this association were also statistically significant.

Table 5 presents the association between increasing serum perfluorooctanoic acid and perfluorooctane sulfonate and the presence of hyperuricemia. We observed a positive association between increasing quartiles of serum perfluorooctanoic acid and perfluorooctane sulfonate and the

odds of hyperuricemia in the unadjusted, age-adjusted, and gender-adjusted models, as well as the multivariable-adjusted models. Models evaluating trend in this association were also statistically significant.

Discussion

In a representative sample of US adults, we found that serum perfluoroalkyl chemical levels were positively associated with elevated uric acid. The association was found to be independent of age, gender, race-ethnicity, education, smoking, alcohol intake, body mass index, hypertension, diabetes, and serum total cholesterol. We found that both perfluorooctanoic acid and perfluorooctane sulfonate levels were associated with elevated uric acid in separate analyses. Our results contribute to the existing literature, which had demonstrated an association between perfluoroalkyl chemicals and uric acid in groups who were exposed to very high perfluoroalkyl chemical levels, by suggesting that perfluoroalkyl chemical levels are positively related to uric acid, even at low exposure levels as seen in the US general population.

Hyperuricemia is the underlying mechanism in gout, a clinical condition where high levels of serum uric acid result in deposition of monosodium urate crystals in the synovial fluid of joints and in the interstitial space and tubules of renal cells resulting, respectively, in arthritis and kidney disease.¹¹ Furthermore, emerging animal and human studies suggest

Table 3 Association between plasma PFOA level and uric acid levels by gender

Quartile	Men			Women		
	Sample size	Unadjusted mean change in uric acid, mg/dL (95% CI)	Multivariate-adjusted mean change in uric acid, mg/dL (95% CI) [‡]	Sample size	Unadjusted mean change in uric acid, mg/dL (95% CI)	Multivariate-adjusted mean change in uric acid, mg/dL (95% CI) [‡]
PFOA*						
Quartile 1	463	1 (referent)	1 (referent)	521	1 (referent)	1 (referent)
Quartile 2	455	0.14 (−0.03, 0.30)	0.08 (−0.06, 0.22)	476	0.16 (0.01, 0.31)	0.14 (−0.03, 0.30)
Quartile 3	471	0.39 (0.18, 0.61)	0.33 (0.15, 0.51)	532	0.37 (0.20, 0.54)	0.29 (0.15, 0.43)
Quartile 4	468	0.48 (0.32, 0.64)	0.41 (0.24, 0.57)	497	0.50 (0.32, 0.69)	0.33 (0.12, 0.54)
P trend		<0.0001	<0.0001		<0.0001	0.0012
Log-transformed PFOA	1857	0.25 (0.13, 0.37)	0.21 (0.09, 0.34)	2026	0.27 (0.18, 0.35)	0.16 (0.06, 0.26)
PFOS[†]						
Quartile 1	466	1 (referent)	1 (referent)	501	1 (referent)	1 (referent)
Quartile 2	460	0.28 (0.06, 0.50)	0.20 (−0.02, 0.43)	509	0.07 (−0.15, 0.29)	0.13 (−0.05, 0.32)
Quartile 3	470	0.26 (0.04, 0.48)	0.20 (−0.01, 0.41)	508	0.32 (0.10, 0.55)	0.19 (−0.04, 0.41)
Quartile 4	461	0.28 (0.06, 0.49)	0.23 (0.02, 0.44)	508	0.42 (0.16, 0.69)	0.23 (−0.02, 0.48)
P trend		0.0223	0.0499		0.0006	0.0706
Log-transformed PFOS		0.16 (0.06, 0.26)	0.11 (0.00, 0.22)		0.24 (0.09, 0.39)	0.13 (−0.01, 0.27)

Notes: *Plasma PFOA quartiles: quartile 1 (<2.4 ppb), quartile 2 (2.4–3.4 ppb), quartile 3 (3.5–5.1 ppb), quartile 4 (>5.1 ppb); in men: quartile 1 (<3.2 ppb), quartile 2 (3.2–4.5 ppb), quartile 3 (4.6–6.4 ppb), quartile 4 (>6.4 ppb); [†]plasma PFOS quartiles: quartile 1 (<11.2 ppb), quartile 2 (11.2–17.8 ppb), quartile 3 (17.9–27.9 ppb), quartile 4 (>27.9 ppb); in men: quartile 1 (<17.5 ppb), quartile 2 (17.5–24.7 ppb), quartile 3 (24.8–35.6 ppb), quartile 4 (>35.6 ppb); [‡]adjusted for age (years), gender (men, women), race-ethnicity (non-Hispanic whites, non-Hispanic blacks, Mexican Americans, others), education categories (<high school, high school, >high school), smoking (never, former, current), alcohol intake (heavy drinker), body mass index (kg/m²), hypertension (absent, present), diabetes (absent, present), and serum total cholesterol (mg/dL). **Abbreviations:** CI, confidence interval; PFOA, perfluorooctanoic acid; PFOS, perfluorooctane sulfonate.

that even mild elevations in serum uric acid may be associated with increased risk of developing hypertension, chronic kidney disease, and cardiovascular outcomes.²² Therefore, identifying environmental factors associated with elevated levels of serum uric acid may have clinical and public health significance.

There is limited previous research available on the effect of perfluoroalkyl chemicals on uric acid, with only two occupational studies and one community-based study examining this hypothesis. In a cross-sectional study of 1025 active workers exposed to perfluorooctanoic acid from a chemical plant based in the US, Sakr et al^{12,13} reported a modest positive association between serum perfluorooctanoic acid levels and uric acid. In a more recent study of 53 male workers at a perfluorooctanoic acid production plant from Italy, Costa et al¹⁴ reported a similar positive association between serum perfluorooctanoic acid levels and uric acid. In a large, community-based, cross-sectional study of perfluorooctanoic acid and perfluorooctane sulfonate and uric acid among 54,951 adult community residents in Ohio and West Virginia, who lived or worked in six water districts contaminated with perfluorooctanoic acid from a chemical plant, Steenland et al¹⁵ reported that the risk of hyperuricemia risk increased modestly with increasing perfluorooctanoic acid; the OR by quintile of perfluorooctanoic acid reported in that study were 1.00, 1.33

(95% CI, 1.24–1.43), 1.35 (95% CI, 1.26–1.45), 1.47 (95% CI, 1.37–1.58), and 1.47 (95% CI, 1.37–1.58; test for trend, $P < 0.0001$). However, all of these studies were based on samples of individuals exposed to perfluorooctanoic acid levels that are much higher than the background exposure levels in the US general population.

Exposure levels of perfluoroalkyl chemicals are important, because a recent study showed that the half-life of perfluorooctanoic acid was 2.9 years at higher exposure levels and 8.5 years at lower exposure levels, suggesting a concentration-dependent clearance of these chemicals from the human body.¹⁶ These half-life values are higher than those reported in previous studies examining occupational workers.^{23,24} Therefore, there is a need to examine the putative association between serum perfluoroalkyl chemical levels and uric acid at these lower levels also. It is in this context that the results of our study employing the nationally representative NHANES survey are relevant. We found that both serum perfluorooctanoic acid and perfluorooctane sulfonate levels are positively associated with uric acid at low exposure levels seen in the US general population, even after adjusting for a variety of confounders.

The biological mechanisms underlying this observed positive association between serum uric acid and perfluoroalkyl chemical levels are not clear. Recent studies have reported that perfluorooctanoic acid exposure may be

Table 4 Association between plasma PFOA level and uric acid levels according to body mass index categories

Quartile	BMI < 30 kg/m ²			BMI ≥ 30 kg/m ²		
	Sample size	Unadjusted mean change in uric acid, mg/dL (95% CI)	Multivariate-adjusted mean change in uric acid, mg/dL (95% CI) [‡]	Sample size	Unadjusted mean change in uric acid, mg/dL (95% CI)	Multivariate-adjusted mean change in uric acid, mg/dL (95% CI) [‡]
PFOA*						
Quartile 1	653	1 (referent)	1 (referent)	331	1 (referent)	1 (referent)
Quartile 2	597	0.17 (0.01, 0.33)	0.16 (0.04, 0.27)	334	0.24 (-0.02, 0.49)	0.11 (-0.11, 0.32)
Quartile 3	665	0.37 (0.19, 0.54)	0.33 (0.20, 0.46)	338	0.52 (0.25, 0.79)	0.46 (0.22, 0.70)
Quartile 4	635	0.52 (0.35, 0.68)	0.40 (0.26, 0.55)	330	0.65 (0.34, 0.96)	0.49 (0.25, 0.73)
P trend		<0.0001	<0.0001		<0.0001	<0.0001
Log-transformed PFOA	2550	0.45 (0.36, 0.55)	0.21 (0.13, 0.30)	1333	0.46 (0.26, 0.66)	0.23 (0.05, 0.41)
PFOS[†]						
Quartile 1	624	1 (referent)	1 (referent)	343	1 (referent)	1 (referent)
Quartile 2	653	0.12 (-0.10, 0.34)	0.22 (0.06, 0.38)	316	0.22 (-0.01, 0.44)	0.11 (-0.08, 0.30)
Quartile 3	652	0.22 (0.06, 0.38)	0.16 (-0.02, 0.34)	326	0.40 (0.05, 0.75)	0.34 (0.01, 0.67)
Quartile 4	621	0.31 (0.12, 0.50)	0.34 (0.18, 0.50)	348	0.28 (-0.06, 0.61)	0.13 (-0.18, 0.44)
P trend		0.0015	0.0005		0.0600	0.2348
Log-transformed PFOS		0.42 (0.35, 0.48)	0.16 (0.08, 0.24)		0.39 (0.18, 0.59)	0.10 (-0.11, 0.30)

Notes: *Plasma PFOA quartiles: quartile 1 (<2.4 ppb), quartile 2 (2.4–3.4 ppb), quartile 3 (3.5–5.1 ppb), quartile 4 (>5.1 ppb); in men: quartile 1 (<3.2 ppb), quartile 2 (3.2–4.5 ppb), quartile 3 (4.6–6.4 ppb), quartile 4 (>6.4 ppb); †plasma PFOS quartiles: quartile 1 (<11.2 ppb), quartile 2 (11.2–17.8 ppb), quartile 3 (17.9–27.9 ppb), quartile 4 (>27.9 ppb); in men: quartile 1 (<17.5 ppb), quartile 2 (17.5–24.7 ppb), quartile 3 (24.8–35.6 ppb), quartile 4 (>35.6 ppb); ‡adjusted for age (years), gender (men, women), race-ethnicity (non-Hispanic whites, non-Hispanic blacks, Mexican Americans, others), education categories (<high school, high school, >high school), smoking (never, former, current), alcohol intake (heavy drinker), body mass index (kg/m²), hypertension (absent, present), diabetes (absent, present), and serum total cholesterol (mg/dL). **Abbreviations:** BMI, body mass index; CI, confidence interval; PFOA, perfluorooctanoic acid; PFOS, perfluorooctane sulfonate.

related to oxidative stress in liver^{25,26} and endothelial cells. Similarly, elevations in serum uric acid may be an indirect measure of oxidative stress.²⁷ Another possibility is that organic anion transporters 1 and 3, which are involved in the tubular secretion of uric acid,²⁸ have been reported to have

a high affinity for perfluorooctanoic acid.²⁹ Therefore, it is possible that perfluorooctanoic acid may compete with the excretion of uric acid.

The main strengths of our study include its population-based nature, inclusion of a representative multiethnic sample,

Table 5 Association between plasma PFOA level and hyperuricemia

Plasma PFC level*	Number at risk (hyperuricemia weighted %)	Unadjusted OR (95% CI)	Age, gender-adjusted OR (95% CI)	Multivariate-adjusted OR (95% CI) [‡]
PFOA*				
Quartile 1	984 (13.6)	1 (referent)	1 (referent)	1 (referent)
Quartile 2	931 (15.7)	1.18 (0.82, 1.69)	1.17 (0.81, 1.69)	1.14 (0.78, 1.67)
Quartile 3	1003 (22.0)	1.79 (1.27, 2.51)	1.84 (1.33, 2.55)	1.90 (1.35, 2.69)
Quartile 4	965 (23.7)	1.97 (1.50, 2.59)	1.97 (1.48, 2.62)	1.97 (1.44, 2.70)
P trend		<0.0001	<0.0001	<0.0001
Log-transformed PFOA		1.62 (1.35, 1.93)	1.42 (1.19, 1.71)	1.43 (1.16, 1.76)
PFOS[†]				
Quartile 1	967 (14.8)	1 (referent)	1 (referent)	1 (referent)
Quartile 2	969 (18.9)	1.34 (1.00, 1.81)	1.38 (1.02, 1.87)	1.46 (1.11, 1.91)
Quartile 3	978 (22.6)	1.68 (1.20, 2.34)	1.63 (1.15, 2.32)	1.69 (1.19, 2.40)
Quartile 4	969 (20.8)	1.51 (1.01, 2.24)	1.42 (0.94, 2.16)	1.48 (0.99, 2.22)
P trend		0.0180	0.0611	0.0433
Log-transformed PFOS		1.47 (1.20, 1.79)	1.21 (0.98, 1.50)	1.21 (0.97, 1.51)

Notes: *Plasma PFOA quartiles: quartile 1 (<2.4 ppb), quartile 2 (2.4–3.4 ppb), quartile 3 (3.5–5.1 ppb), quartile 4 (>5.1 ppb); in men: quartile 1 (<3.2 ppb), quartile 2 (3.2–4.5 ppb), quartile 3 (4.6–6.4 ppb), quartile 4 (>6.4 ppb); †plasma PFOS quartiles: quartile 1 (<11.2 ppb), quartile 2 (11.2–17.8 ppb), quartile 3 (17.9–27.9 ppb), quartile 4 (>27.9 ppb); in men: quartile 1 (<17.5 ppb), quartile 2 (17.5–24.7 ppb), quartile 3 (24.8–35.6 ppb), quartile 4 (>35.6 ppb); ‡adjusted for age (years), gender (men, women), race-ethnicity (non-Hispanic whites, non-Hispanic blacks, Mexican Americans, others), education categories (<high school, high school, >high school), smoking (never, former, current), alcohol intake (heavy drinker), body mass index (kg/m²), hypertension (absent, present), diabetes (absent, present), and serum total cholesterol (mg/dL). **Abbreviations:** CI, confidence interval; OR, odds ratio; PFOA, perfluorooctanoic acid; PFOS, perfluorooctane sulfonate; PFC, perfluoroalkyl chemicals.

adequate sample size, objective measurement of diabetes status following current guidelines, and the availability of extensive data on confounders for multivariate adjustment. The main limitation of our study is the cross-sectional nature of NHANES. Therefore, similar to previous studies that examined the association between other environmental exposures and disease states using the NHANES data, the temporal nature of the association between perfluoroalkyl chemicals and diabetes cannot be concluded from the current study. Another limitation is that adjustment for variables that are potential mediators, such as hypertension in the multivariate model, may be an overadjustment, and therefore may have led to an underestimation of the true association between serum perfluoroalkyl chemical levels and uric acid. It is possible that the true association may be in fact of stronger magnitude.

In summary, we found that serum perfluoroalkyl chemical levels were positively associated with elevated uric acid in a representative, multiethnic sample of US adults. This association was found to be independent of age, gender, race-ethnicity, education, smoking, alcohol intake, body mass index, hypertension, diabetes, and serum total cholesterol. Our results contribute to the literature by suggesting that perfluoroalkyl chemical levels are related to uric acid, even at low exposure levels, in the general population.

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Disclosure

The authors report no conflicts of interest in this work.

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