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Exposed Population of Community Residents and Workers in
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Ulcerative Colitis and Perfluorooctanoic Acid (PFOA) in a Highly Exposed Population of Community Residents and Workers in the Mid-Ohio Valley

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Abbreviations

BMI	Body mass index
CI	Confidence interval
EPA	Environmental Protection Agency
IL3	interleukin 3
PFOA	Perfluorooctanoic acid
PPAR	peroxisome proliferator-activated receptors
ng/ml	nanogram/milliliter

Abstract

Background: Little is known about environmental determinants of autoimmune diseases.

Objectives: To study autoimmune disease in relation to perfluorooctanoic acid (PFOA), which was introduced in the late 1940s and is now ubiquitous in the serum of residents of industrialized countries.

Methods: In 2008-2011 we interviewed 32,254 U.S. adults with high serum PFOA serum levels (median 28 ng/ml) from drinking contaminated water near a chemical plant. Disease history was assessed retrospectively from 1952 or birth (if later than 1952) until interview. Self-reported history of autoimmune disease was validated via medical records. Cumulative exposure to PFOA was derived from estimates of annual mean serum PFOA levels during follow up, which were based on plant emissions, residential/work history, and a fate-transport model. Cox regression models were used to estimate associations between quartiles of cumulative PFOA serum levels and the incidence of autoimmune diseases with 50 or more validated cases, including ulcerative colitis (n=151), Crohn's disease (n=96), rheumatoid arthritis (n=346), insulin-dependent-diabetes (presumed to be Type 1) (n=160), lupus (n=75), and multiple sclerosis (n=98).

Results: The incidence of ulcerative colitis (UC) was significantly increased in association with PFOA exposure, with adjusted rate ratios by quartile of exposure of 1.00, 1.76 (95%CI: 1.04, 2.99), 2.63 (95% CI: 1.56, 4.43), and 2.86 (95% CI: 1.65, 4.96) (trend p-value < 0.0001). A prospective analysis of ulcerative colitis diagnosed after the baseline 2005-2006 survey (n = 29 cases) suggested a positive but non-monotonic trend (test for trend, p=0.21).

Discussion: To our knowledge, this is the first study of associations between this common environmental exposure and autoimmune diseases in humans. We found evidence that PFOA is associated with ulcerative colitis.

Introduction

Autoimmune diseases are estimated to affect about 7% of the US population (Miller et al. 2012, Cooper et al. 2009). The most common include inflammatory bowel disease (consisting of ulcerative colitis and Crohn's disease), rheumatoid arthritis, celiac disease, multiple sclerosis, systemic sclerosis, lupus (systemic lupus erythematosus), Type 1 diabetes, and two types of thyroid disease (Grave's disease, Hashimoto's thyroiditis)(Cooper et al. 2009). In general these diseases are chronic and incurable. They are thought to occur due to an interaction between genes and the environment. Twin studies on concordance suggest a relatively low genetic contribution for rheumatoid arthritis, ulcerative colitis, and systemic sclerosis, a modest contribution for Crohn's disease and lupus, and a high genetic contribution for Type 1 diabetes and auto-immune thyroid disease. Data are largely lacking for other autoimmune diseases (Bogdanos et al. 2012, Baumgart and Carding 2007). Genetic studies, including recent genome-wide association studies (GWAS), have identified a number of genes that may contribute to autoimmune diseases, particularly genes of the major histocompatibility complex (MHC) that have associated with rheumatoid arthritis, systemic sclerosis, and lupus (Delgado-Vega et al. 2012). A number of specific genes have been associated with more than one autoimmune disease.

To date few environmental risk factors for autoimmune disease have been identified. The principal ones are 1) crystalline silica, which has been associated with rheumatoid arthritis, systemic sclerosis, and lupus, 2) smoking, positively associated with seropositive rheumatoid arthritis, and negatively associated with ulcerative colitis, 3) solvents associated with systemic sclerosis, and 4) sunlight, which has been negatively associated with multiple sclerosis (Miller et al. 2009). Although information on population trends in autoimmune disease rates is limited,

concerns have been raised that autoimmune disease incidence rates may be increasing due to increasing exposures to xenobiotics (Miller et al. 2012). One such xenobiotic, which is ubiquitous in the serum of US residents, and which has been shown to affect immune responses in rodents (Dewitt et al. 2009), is perfluorooctanoic acid (PFOA). To our knowledge, there are no prior studies of associations between PFOA and autoimmune disease in humans.

Here we focus on a large cohort (n=32,254) that has been exposed to high levels of PFOA, a perfluorinated compound introduced into the environment in the 1950s. PFOA (also known as C8) is ubiquitous at low levels (ng/ml) in the serum of virtually all residents of industrialized countries (Lindstrom et al. 2011). Human exposure to PFOA occurs via many sources, including food, drinking water (Shin et al. 2011a; Shin et al. 2011b), house dust (Strynar and Lindstrom 2008), and air (Fromme et al. 2009). PFOA has been used until recently in manufacturing a wide variety of consumer products, such as Gore-tex™, Teflon™, and Scotchgard™ (Shin et al. 2011a). The cohort studied here was exposed primarily due to drinking water contaminated with PFOA that originated from emissions from a nearby chemical plant that used PFOA to manufacture Teflon™. Median PFOA serum levels in this population in 2005-2006 were 28 ng/ml, and mean levels were 72 ng/ml, compared with an estimated mean serum concentration of 4 ng/ml for the general US population (Steenland et al. 2009; Calafat et al. 2007).

PFOA is not biodegradable and persists indefinitely in the environment. In humans it is thought to be stored in the liver, kidney, and serum, and has an elimination half-life of approximately 3.5 years (Olsen et al. 2007). PFOA is a rodent carcinogen and causes fetal loss in mice at dietary levels of approximately 20 ppm (Lau et al. 2007). Rodent studies have reported evidence of immunosuppressant effects including lymphoid organ atrophy and decreased de novo antibody production in certain strains of mice (DeWitt et al. 2008; Yang et al. 2000). To date information

about PFOA in relation to immune function in humans is limited to a few cross-sectional studies of relatively insensitive immune markers (as reviewed in Steenland et al. 2010). No clear patterns of immune suppression have been reported based on these studies. However, a recent study of children in the Faroe Islands reported a significant decrease in childhood antibody levels to diphtheria and tetanus at age 7 in association with serum levels of PFOA and another fluorocarbon, PFOS (perfluorooctane sulfonic acid)(Grandjean et al. 2012). To our knowledge, there are no published data on autoimmune disease in relation to PFOA in animals or humans.

The autoimmune diseases considered here are ulcerative colitis, Crohn's disease, rheumatoid arthritis, multiple sclerosis, lupus, and Type 1 diabetes. For these six diseases we had data on self-reported occurrence, for which we then sought validation through medical records review. Analyses here are limited to cases validated based on medical records data. For most other autoimmune diseases we did not have self-reported data, or the numbers of cases were too small to analyze. Self-reported cases of Graves' disease and Hashimoto's thyroiditis are not included in this analysis because they were difficult to distinguish from other (non-autoimmune) forms of hyperthyroidism and hypothyroidism.

While it is likely that different autoimmune diseases have separate etiologies, the fact that several genes have been linked to more than one autoimmune disease also suggests that they may have some aspects in common (Delgado-Vega et al. 2012). Here we have analyzed the relationship of each autoimmune disease to PFOA.

This study is one of a series of studies conducted by the C8 Science Panel, a three-person panel of epidemiologists set up pursuant to a 2004 legal settlement between plaintiffs and DuPont (see www.c8sciencepanel.org, and Frisbee et al. 2009). The settlement mandated a baseline survey of

69,000 individuals who lived or who had lived in six water districts where water had been contaminated with PFOA (called the C8 Health Project). The settlement also created the C8 Science Panel, which was charged with determining whether it is more probable than not that PFOA is ‘linked to’ (associated with) any human disease. The C8 Science Panel conducted eleven studies over the course of 5 years to make this determination.

Methods

Study population and exposure estimation

Community cohort

We studied a mid-Ohio valley community-based population that comprised individuals who lived or worked in any of six PFOA contaminated water districts and participated in the C8 Health Project baseline survey in 2005-2006 (Frisbee et al. 2009). The principal route of exposure for this population was via drinking water contaminated with PFOA. Approximately 80% of current (2005-2006) residents in the six districts participated in the C8 Health Project (Frisbee et al. 2009).

C8 Health Project participants (n=69,030) had their PFOA serum levels measured and provided a medical history. Most adult C8 Health Project participants (74% of participants age 20 or above) consented to participate in a follow-up study. Of these we subsequently interviewed 82% (n = 32,712). These interviews form the basis for the current study. For further details of the interview process, see Winqvist et al. (2013)

Two rounds of surveys were conducted, the first from August 2008 to April 2010 and the second from May 2010 to May 2011. Surveys were completed over the phone (63%) or online (37%), and most participants completed both rounds of surveys. Both surveys included the same

questions regarding demographic information and medical history, including questions about whether they ever were told by a doctor that they had specific chronic diseases. For participants who died after completing the C8 Health Project survey in 2005/2006, or who were not capable of completing a follow-up survey we surveyed their next-of-kin (4% of the cohort had next-of-kin interview). The Emory IRB reviewed and approved all aspects of this study, including consent forms and surveys.

To estimate past exposures, we developed historical yearly serum PFOA estimates for community residents based on the estimated intake of PFOA-contaminated drinking water, assuming low background exposure. The estimates of drinking water PFOA concentrations were based on the amount of PFOA released from the DuPont plant, wind patterns, river flow, groundwater flow, and the residential address history provided by each participant (Shin et al., 2011a, b). Each estimated yearly serum concentration took into account new exposure during the year and the estimated amount of PFOA remaining in the body after partial excretion during the prior year. The final community cohort comprised 28,541 community residents with estimated historical serum concentrations who had never worked at the DuPont plant.

Workers cohort

In addition to the community cohort, we studied a cohort of 4,391 past and current workers at the DuPont plant using the same survey completed by the community cohort participants, including a residential history. This group was a subset of a cohort of 6,027 workers employed at the DuPont chemical plant between 1948 and 2002 who were previously studied for mortality (Leonard et al. 2009; Steenland et al. 2012).

Estimates of annual serum PFOA levels for workers in different jobs were developed by the C8 Science Panel (Woskie et al. 2012) and combined with estimated annual serum levels from residential exposure to contaminated drinking water that were derived as described above for the community cohort. We estimated combined residential and occupational exposure for 3,713 (84%) workers who completed a follow up survey, including 1,890 (51%) who also had participated in the 2005/2006 C8 Health Project.

Combined Community and Worker Population

Community residents and workers were combined to form a final population of 32,254 people for whom we could study the relationship between past PFOA serum levels and subsequent disease. Our model-based exposure predictions were well correlated with serum PFOA measurements obtained in 2005-2006 for community cohort members ($r=0.67$)(Shin et al. 2011a).

Medical history

All participants were asked about lifetime medical history, focusing on chronic disease. Participants who reported that they had been told they had one of the specific chronic diseases of interest were then asked the age at first diagnosis, and for consent to review medical records, as well as the relevant health care provider. Approximately 60% of the cohort reported having had a disease of interest; 79% percent of these consented for us to review their medical history. Professional medical abstractors obtained records documenting the disease in question from medical providers by mail or by visiting their office. Overall, we were able to find a medical record relevant to the reported disease for 92% of those consenting to medical record review (Winquist et al. 2013). For three diseases, multiple sclerosis, rheumatoid arthritis, and lupus, we

were also able to use supplemental data from the C8 Health Project (2005-2006), which conducted its own medical record validation, to confirm some cases that had been self-reported to us, but for whom we were unable to obtain a medical record.

Autoimmune disease in the cohort

Participants were asked whether a doctor or other health professional had ever told them they had an autoimmune disease, with specific categories listed for lupus, multiple sclerosis, myasthenia gravis, Sjögren's syndrome, vasculitis, Addison's disease, or other. If the answer was 'other', the participant was asked to specify the disease. In a separate question, subjects were asked whether a doctor or health professional had ever told them they had had inflammatory bowel disease, excluding irritable bowel syndrome. They were then asked to specify whether the inflammatory bowel disease was ulcerative colitis or Crohn's disease. In another question, participants were asked whether a doctor had ever told them they had arthritis, and if yes, whether it was rheumatoid or osteoarthritis. Finally, participants were asked if they had ever had diabetes, with specific categories of Type 1, or Type II (excluding pregnancy-induced diabetes).

Analyses

Associations between PFOA exposure and each outcome were conducted using separate survival analyses (Cox regression) with age as the time variable. Follow-up began in 1952 (approximate date of first emissions of PFOA from the DuPont plant), or date of birth, whichever came later. Years outside the study area were included in follow-up and assigned US background levels, although sensitivity analyses were conducted excluding person-time prior to arrival in the mid-Ohio valley and/or excluding background levels in calculating cumulative exposure. For each

outcome, follow-up ended at time of last interview, at time of disease occurrence, or at the time of death, whichever came earlier. Cases of each outcome were restricted to cases validated based on medical record review. Participants who self-reported an outcome that was not validated were excluded from the analysis for that outcome.

We estimated associations with cumulative exposure to PFOA, which was calculated by summing estimated yearly serum concentrations during follow-up and modeled as a time-dependent variable in Cox regression. We also considered cumulative exposure with a lag of 10 years (i.e., discounting exposure in the prior 10 years). We conducted analyses by quartile of cumulative exposure, with cutpoints for each outcome determined from the cumulative exposure of the cases of that outcome at time of diagnosis, and separate cutpoints derived for lagged and unlagged exposures. Rate ratios (the ratio of disease rates) were estimated for the 2nd, 3rd, and 4th quartiles relative to the first quartile, which was the referent group. Tests for trend were based on the p-value of the coefficient of the natural log of cumulative exposure modeled as a continuous time-dependent variable in a Cox regression model that included the same covariates as the corresponding model of cumulative PFOA exposure as a categorical variable.

All models were adjusted for a standard set of covariates including gender and race/ethnicity (white or non-white) and time-dependent variables for smoking (current, former, vs. never), BMI (body mass index) (<18.5, 18.5-24.9, 25-29.9, ≥ 30.0 kg/m²), and alcohol consumption (current, former, vs. never) based on the most recent survey. The proportional hazard assumption of a constant hazard ratio for cumulative exposure over time was verified based on the non-significance of an interaction term between time (age) and cumulative exposure.

We conducted two types of sensitivity analyses. In one ('qualifying time analysis'), the time of follow-up started when the participant qualified for the C8 Health Project (i.e., time of first residence in a PFOA-contaminated water district, or first employment at the plant). This eliminated prior non-exposed person-time before cohort eligibility, and resulted in a loss of about 10%-15% of cases depending on the outcome being evaluated. Second, we estimated associations with cumulative exposures above background levels only, so that exposure began at year of first exposure to contaminated water or work in the DuPont plant ('above background' analysis). These analyses included the same number of participants for each outcome as corresponding models of associations with cumulative PFOA exposure including estimated background levels of exposure (approximately 4 ng/ml serum).

Other analyses were restricted to person-time after the C8 Health Project was established in 2005-2006 (prospective analysis), with participants reporting a history of any of the autoimmune diseases before participation in the C8 Health Project (prevalent cases) excluded to restrict the analysis to incident cases diagnosed after 2005-2006. Person-time for participants in the C8 Health Project started on the date of their C8 Health Project blood draw and interview, which varied from July 2005-July 2006. For members of the worker cohort who did not participate in the C8 Health Project (n = 1,823), person-time at risk began on July 1, 2006. For all but two of the autoimmune diseases (rheumatoid arthritis and ulcerative colitis) there were insufficient cases to conduct a prospective analysis.

Results

Table 1 provides descriptive statistics for the cohort. The median year of birth of the cohort was 1957, 54% were female, 20% were current smokers, 27% were current drinkers, 18% had a college education or more, and 36% were obese (BMI >30) (BMI, smoking, and drinking

assessed at time of last interview). PFOA serum levels in 2005-2006 were 24 ng/m³ for community residents and 113 ng/m³ for workers. The median length of follow up was 53 years (birth to end of follow-up); the median length of follow-up after the earliest year living or working in a contaminated water district was 29 years.

Numbers of validated cases of each autoimmune disease are shown along with mean at diagnosis and gender in Table 2. Validated cases (based on medical record review) included 346 participants who reported rheumatoid arthritis for which they were taking medication (25% of the 1,292 who reported it, 72 participants with lupus (39% of 187 self-reported cases), and 98 with multiple sclerosis (65% of 150 self-reported cases). There were 151 validated cases of ulcerative colitis (25% of self-reported cases), and 96 validated cases of Crohn's disease (53% of self-reported cases) (ten cases were validated for both ulcerative colitis and Crohn's disease).

Of the 342 self-reported cases of Type 1 diabetes, 69 were validated as either Type 1 diabetes or insulin-dependent diabetes (Type 1 diabetes is also known as insulin-dependent diabetes), and we found another 91 validated cases (as either insulin-dependent or Type 1) among subjects who reported that they had diabetes but did not know what kind. Of the 160 validated cases, 85 were validated specifically for Type 1 diabetes, while the remaining cases were validated as 'insulin-dependent'. We conducted analyses for both the group of 160 validated cases and also for the sub-group of 85 cases specifically validated as Type 1 diabetes.

The known higher incidence of rheumatoid arthritis, lupus, and multiple sclerosis among women compared with men is reflected in our data (Table 2). The age at diagnoses also tend to conform to what is known, e.g., most of these diseases tend to occur during middle age. The mean age for Type 1 diabetes (combining insulin-dependent and specifically Type 1) is younger than the mean

age at onset for the other diseases reflecting the fact that some of these cases occur among juveniles, although in recent years the majority of Type 1 diabetes occurs among adults.

Table 3 shows the results of the Cox regression for the overall analysis. Ulcerative colitis showed a significant positive association with cumulative PFOA exposure, with monotonically increasing rate ratios for both the unlagged and lagged exposures and significant trends based on models of log-transformed cumulative exposure. No such trend was evident for Crohn's disease, or any of the other autoimmune diseases examined.

Sensitivity analyses using either exposures beginning in the 'qualifying year' or without background added in, were similar to those shown in Table 3 (data not shown). For ulcerative colitis, with no background exposures added, the quartile RRs were 1.00, 1.27 (95% CI: 0.78, 2.08), 2.08 (95% CI: 1.26, 3.44), and 2.30 (95% CI: 1.36, 3.91) (test for trend $p < 0.0001$). The corresponding RRs with estimated person-time before moving to an exposed water district were 1.59 (95% CI: 0.95, 2.65), 2.40 (95% CI: 1.44, 3.99), and 2.33 (95% CI: 1.37, 3.97) (test for trend $p < 0.0001$). Again, ulcerative colitis was the only outcome associated with PFOA.

Only ulcerative colitis ($n = 30$) and rheumatoid arthritis ($n = 56$) had ≥ 25 cases diagnosed after participation in the original C8 Health Project (2005-2006) or July 1 2006 (for those enrolled in the worker cohort who did not participate in the C8 Health Project). All RRs for ulcerative colitis were elevated after the first quartile, but without a monotonic trend, based on this prospective analysis (trend $p = 0.21$, unlagged analysis) (Table 4).

Discussion

For all six autoimmune diseases considered here, the prevalence in the Mid-Ohio Valley conformed broadly with that expected from US and Western European populations, although

expected prevalences for some of these diseases are uncertain (Cooper et al. 2009). Crude prevalences for validated cases per 100,000 adults (≥ 20 years of age) in our cohort in 2008-2001 were 410, 300, 1080, 500, 230, and 310 for ulcerative colitis, Crohn's disease, rheumatoid arthritis, Type 1 diabetes (including insulin-dependent), lupus, and multiple sclerosis, respectively. Data from US and Western Europe based on both hospital and non-hospital data suggest comparable patterns but lower prevalences for adults and children combined, with ranges of 140-300, 100-200, 310-830, 340-570, 30-150, and 50-360, respectively, for these same diseases (Cooper et al. 2009). On the other hand a recent study based on medical claims data (n=12 million) reported prevalences of 263 and 241 per 100,000 US adults for ulcerative colitis and Crohn's disease, respectively, which suggests that ulcerative colitis may be in excess in our population (Kappleman et al. 2012). These data also indicate that the prevalences of both types of adult IBD have been slightly increasing over the 6 year period studied (2004-2009).

Cumulative PFOA exposures were associated with ulcerative colitis, but not Crohn's disease, in our study population. Ulcerative colitis and Crohn's disease are clinically distinct conditions, with distinguishing clinical, anatomical, and histological findings. Crohn's disease occurs in any part of the gastrointestinal tract, although a majority of the cases start in the end of the small bowel. Ulcerative colitis is restricted to the colon and the rectum. Ulcerative colitis affects the mucosa (epithelial lining of the gut), while Crohn's disease affects the whole bowel wall. However, there is no definitive diagnostic method to distinguish between them in about 10% of cases (Hanauer 2006; Sands 2004).

Smoking is positively associated with Crohn's disease (Mahid et al. 2006). In contrast, current smoking is negatively associated with ulcerative colitis, though former smoking is a risk factor (Mahid et al. 2012). A meta-analysis for smoking and ulcerative colitis found an RR= 0.58 for

current smoking, and RR= 1.76 for former smoking (Mahid et al. 2006). It is reassuring that we found some support for these patterns in our data. Specifically, RRs for current and former smoking and ulcerative colitis were 0.63 (95% CI: 0.39-1.01; p=0.05) and 1.90 (95% CI 1.29-2.82, p=0.001), respectively. The RRs for current and former smokers for Crohn's disease were 1.26 (95% CI: 0.28-2.03; p=0.35) and 0.94 (95% CI: 0.50-1.76; p=0.85), respectively.

Both ulcerative colitis and Crohn's disease have a genetic component, but twin studies suggest this component is relatively modest for ulcerative colitis. The concordance of ulcerative colitis in monozygotic twins is 10%, and 3% in dizygotic twins, compared with 37% and 7%, respectively, for Crohn's disease (Baumgart and Carding 2007). Ordas et al. (2012) have recently reviewed environmental risk factors for ulcerative colitis. Positive risk factors include prior gastrointestinal infections and oral contraceptive use, while protective factors include appendectomy and breastfeeding. We do not have data to control for these factors in our analysis. However, with regard to oral contraceptives and breastfeeding, we analyzed our data separately for men and women, and found little difference between these analyses (data not shown).

A growing body of evidence suggests inflammatory bowel disease may be due, in part, to deficiencies in innate immunity and an impaired intestinal epithelial barrier, resulting in damaging inflammatory responses to the microbial environment of the gastrointestinal tract (Baumgart et al. 2007; Ordas et al. 2012; Gersemann et al. 2012). Animal models and human studies suggest that PFOA toxicity includes systemic suppression of adaptive immunity and antibody production (DeWitt 2012; Grandjean 2012), and the alteration of inflammatory pathways (De Witt 2012). Many PFOA-related immune-effects may be mediated through activation of peroxisome proliferator-activated receptors (PPAR)- α , PPAR- γ or other receptors

expressed in the colon, but non-receptor mediated changes may also occur (DeWitt et al. 2012). Although PFOA-related activation of human PPAR- γ has not been established (Takacs 2007; Vanden Heuvel 2006), activation of PPAR- γ through endogenous and exogenous ligands appears to have anti-inflammatory effects in colitis models and is being explored for therapeutic potential (Dubuquoy 2006). These findings seem to contradict our finding of a positive association between PFOA and ulcerative colitis. However, specific gastrointestinal and local immune effects of PFOA are not described in the literature.

Plausible mechanisms linking PFOA and ulcerative colitis may include shifts in the balance of tissue macrophages towards an anti-inflammatory M2 phenotype and/or a TH2-like response to specific antigens (DeWitt et al. 2012). Experimental findings also suggest PPAR-gamma activation may lead to reprogramming of tissue macrophages towards an M2 anti-inflammatory phenotype, which may contribute to decreased vaccine efficacy or immunosuppression in diseases dependent on cytotoxic T-cell responses (DeWitt et al. 2012). Other studies suggest PFOA may induce shifts in the TH1/TH2 balance, increasing production of TH2-like cytokines involved in hypersensitivity responses (Dewitt et al. 2012). One TH2-type cytokine (IL-13) is thought to play a unique and critical role in ulcerative colitis in gut mucosal inflammatory response (Mannon 2012), but PFOA-related effects on IL-13 expression have not been described. Ultimately, the context of the microbial environment may be especially relevant in explaining PFOA-related effects on ulcerative colitis. One study's findings that PFOA fed to mice (0.02% of total diet by weight) leads to both systemic neutropenia and increased lipopolysaccharide-related cytokine release in other cell populations (e.g., macrophages)(Qazi et al. 2009), suggests a general increase in host susceptibility to infections (DeWitt et al. 2012) Together, these findings suggest that the association between PFOA and ulcerative colitis in the present study (in

contrast with a lack of clear associations with Crohn's or any of the other autoimmune diseases examined), might be explained by effects of PFOA on responses to bacterial exposures and other unique aspects of lower gastrointestinal toxicity that may not be reflected by systemic or other organ-specific immune effects.

It is possible that increased gut absorption of PFOA in subjects with ulcerative colitis (including possibly early still undiagnosed ulcerative colitis) could lead to higher measured serum PFOA levels, which could create a false appearance of a causal association, if measured levels were used to predict current (or even subsequent) ulcerative colitis. However, in our data we estimated associations with cumulative PFOA exposure predicted by a model that did not incorporate measured PFOA concentrations, hence 'reverse causality' (by which disease might increase measured serum levels rather than serum levels preceding disease) is not relevant here.

There are a number of limitations to our analysis. Our cohort is largely a survivor cohort, with community members having to have been alive in 2005/2006 at the time of a baseline survey. If community residents with autoimmune disease were less likely to have survived until 2005/2006, they would have been less likely than other residents to have enrolled in the study, possibly biasing our results. However, two factors argue against this as an explanation for our positive findings for ulcerative colitis. First, life expectancy for those with ulcerative colitis is about the same as the general population (Winther et al. 2003; Davoli et al. 1997). Second, our prospective analyses of ulcerative colitis, although limited by small numbers, would be unaffected by this problem, and shows elevated RRs (ranging from 1.6 to 2.1) for all three upper quartiles vs. the lowest quartile of cumulative exposure, although these data are limited by small numbers, and there is no significant trend.

Other limitations include possible inaccuracies in disease validation and ascertainment, including under ascertainment of cases that were not self-reported by participants, lack of validation of the absence of disease among those who did not report autoimmune disease outcomes, and the possible exclusion of cases that could not be verified due to lack of consent or inability to obtain appropriate medical records. We also lacked data on other potential environmental toxicants in the serum of mid-Ohio valley residents that could have confounded associations with PFOA, and given the limited information on the causes of autoimmune diseases, we cannot rule out possible confounding by other unmeasured exposures or characteristics.

Another limitation is that exposures were classified according to estimated historical serum levels based on predicted water concentrations and predicted air concentrations using an environmental fate and transport model, individual residential histories, and maps of public water supply networks, coupled with a one-compartment absorption and excretion model (Shin et al. 2011a). Model-estimated levels were correlated with measured levels in 2005-2006 ($r_s=0.67$), based on 45,000 participants.

There are also a number of strengths to our analysis. We studied a well-defined cohort with documented high exposure to a chemical that has been reported to suppress the immune system in animals. Disease outcomes were confirmed through medical records review, with relatively large numbers for many autoimmune diseases. Nonetheless, our finding of a positive association between cumulative serum PFOA concentration and incident ulcerative colitis, and a lack of associations with other autoimmune diseases, are based only on one study and need replication.

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Table 1. Cohort characteristics based on the most recent survey and measured serum PFOA levels in 2005-2006

Characteristic	Community Cohort (n=28,541)	Worker Cohort (n= 3,713)	Combined Cohorts (n=32,254)
Year of Birth			
25 th percentile	1947	1941	1946
Median	1958	1951	1957
75 th percentile	1970	1963	1969
Gender			
Female	16,602 (58%)	758 (20%)	17,360 (54%)
Race/ethnicity			
White, non-Hispanic	27,901 (98%)	3,284 (88%)	31,185 (97%)
Other	640 (2%)	134 (4%)	774 (2%)
Missing	0	295 (8%)	295 (1%)
Body Mass Index (BMI)			
<18.5 kg/m ³	414 (2%)	38 (1%)	452 (1%)
18.5-<25.0 kg/m ³	7,693 (27%)	972 (26%)	8,665 (27%)
25.0-<30.0 kg/m ³	9,689 (34%)	1,618 (44%)	11,307 (35%)
≥30.0 kg/m ³	10,694 (37%)	1,057 (29%)	11,751 (37%)
Education			
<High School	3,026 (11%)	37 (1%)	3,063 (10%)
High School	11,706 (41%)	1,265 (34%)	12,971 (40%)
Some College	9,441 (33%)	1,081 (29%)	10,522 (33%)
College Diploma or Higher	4,366 (15%)	1,328 (36%)	5,694 (18%)
Missing	2 (<1%)	2 (<1%)	4 (<1%)
Smoking			
Never Smoked	13,527 (47%)	1,989 (54%)	15,516 (48%)
Smoked and quit	8,899 (31%)	1,297 (35%)	10,196 (32%)
Smoked, did not quit	6,115 (21%)	427 (12%)	6,542 (20%)
Regular Alcohol Consumption			
Never	17,011 (60%)	1,683 (45%)	18,694 (58%)
Yes and quit	4,105 (14%)	535 (14%)	4,640 (14%)
Yes, did not quit	7,360 (26%)	1,486 (40%)	8,846 (27%)
In Community Cohort	28,541 (100%)	1,890 (51%)	30,431 (94%)
Serum PFOA Concentration Measurement from the C8 Health Project (2005-2006) (ng/ml) ^a			
Mean	71	325	87
Standard Deviation	151	921	278
Inter-quartile range	12-59	56-256	13-68
Median	24	113	26

^a Serum PFOA data for members of the worker cohort are restricted to 1,890 workers who had serum PFOA measurements because of participation in the C8 Health Project in 2005-2006 . All other data for workers comes from survey data and refers to all 3,713 workers.

Table 2. Numbers of participants with validated autoimmune diseases, mean age at diagnosis, and sex^a

Autoimmune disease	Number self-reported ^b	Number validated ^b	Mean age	% Female ^c
Ulcerative colitis	596	151	44	59
Crohn's disease	178	95	39	59
Rheumatoid arthritis	1292	346	48	69
Type 1 diabetes -broad ^d	342	160	35	52
Type 1 diabetes-narrow ^d	n.a.	85	27	48
Lupus	187	72	43	87
Multiple sclerosis	150	99	38	77

^aCases confirmed through medical records review.

^bIn the entire cohort, among people reporting diseases for which validation was sought, approximately 75% consented to medical record review, and among those who consented, a record was obtained for 92%; among these the percent validated across all diseases was 77%. For multiple sclerosis, rheumatoid arthritis, and lupus we included among the validated those cases which had been self-reported and not confirmed by us for lack of a medical record, but which had been confirmed previously during the C8 Health Project (29, 19, and 5 cases respectively). For rheumatoid arthritis, numbers are restricted to cases who reported taking medication; these were the only ones for which we sought medical records

^c 54% of the entire cohort was female.

^d Type 1 diabetes-broad includes cases self-reported as Type 1 diabetes and then verified as either Type 1 diabetes or insulin- dependent diabetes, while Type 1 diabetes-narrow is restricted to the same self-reported cases specifically validated for Type 1 diabetes.

Table 3. Overall retrospective survival analysis results based on follow-up from 1952 through 2008-2011^a

	Rate ratio quartile 2 vs. 1 (95% CI) ^b	Rate ratio quartile 3 vs. 1 (95% CI)	Rate ratio quartile 4 vs. 1 (95% CI)	Test for trend of log cumulative exposure
Unlagged exposure				
Ulcerative colitis	1.76 (1.04, 2.99)	2.63 (1.56, 4.43)	2.86 (1.65, 4.96)	<0.0001
Crohn's disease	1.25 (0.61, 2.58)	1.15 (0.55, 2.41)	1.00 (0.48, 2.09)	0.73
Rheumatoid arthritis	1.24 (0.85, 1.79)	1.40 (0.96, 2.03)	0.99 (0.68, 1.43)	0.84
Type 1 diabetes- broad ^c	0.68 (0.29, 1.58)	0.53 (0.22, 1.30)	0.54 (0.22, 1.33)	0.84
Type 1 diabetes-narrow	0.83 (0.25, 2.78)	1.41 (0.40, 4.95)	0.88 (0.25, 3.06)	0.68
Lupus	1.49 (0.68, 3.34)	1.01 (0.44, 2.30)	0.71 (0.31, 1.65)	0.94
Multiple sclerosis	0.85 (0.44, 1.63)	1.56 (0.81, 3.00)	1.26 (0.65, 2.42)	0.22
10 year lagged exposure				
Ulcerative colitis	1.71 (0.89, 3.27)	2.05 (1.07, 3.91)	3.05 (1.56, 5.96)	<0.0001
Crohn's disease	0.80 (0.32, 1.99)	0.97 (0.36, 2.60)	0.69 (0.26, 1.82)	0.79
Rheumatoid arthritis	1.53 (0.61, 2.58)	1.73 (1.10, 2.71)	1.35 (0.87, 2.11)	0.73
Type 1 diabetes-broad	0.42 (0.09, 2.00)	0.70 (0.14, 0.35)	0.38 (0.08, 1.93)	0.84
Type 1 diabetes-narrow	0.50 (0.05, 4.90)	1.32 (0.14, 12.40)	0.71 (0.07, 7.14)	0.65
Lupus	0.79 (0.27, 2.34)	1.26 (0.40, 4.03)	0.61 (0.19, 1.91)	0.93
Multiple sclerosis	1.16 (0.54, 2.47)	1.62 (0.75, 3.52)	1.32 (0.61, 2.84)	0.59

^a Follow up continued until the date of the last survey, the date of diagnosis for each individual outcome being evaluated, or on the date of death, whichever came first.

^b Cutpoints for these quartiles varied by disease analyzed, as they were based on dividing the cases equally by quartiles. As an example, for ulcerative colitis, the cutpoints were 158, 586, and 3500 ng/ml years, which for a 50 year old would correspond to average exposure of approximately < 3, 3-11, 11- 70, and 70+ ng/mls per year.

^c Type 1 diabetes-broad includes self-reported cases listed as verified as either Type 1 diabetes or insulin-dependent diabetes, while Type 1 diabetes-narrow is restricted to those self-reported cases specifically validated for Type 1 diabetes.

Table 4. Prospective survival analysis results, follow-up 2005-2006 through 2008-2011, for selected outcomes (≥ 30 cases)

	Rate ratio quartile 2 vs. 1 (95% CI)	Rate ratio quartile 3 vs. 1 (95% CI)	Rate ratio quartile 4 vs. 1 (95% CI)	Test for trend of log cumulative exposure
Unlagged				
Ulcerative colitis	1.68 (0.62, 4.66)	2.10 (0.75, 5.85)	1.62 (0.57, 4.61)	0.21
Rheumatoid	0.81 (0.39, 1.68)	2.07 (1.00, 4.27)	0.52 (0.25, 1.09)	0.89
10 year lag				
Ulcerative colitis	1.21 (0.43, 3.34)	2.16 (0.80, 5.81)	1.51 (0.43, 4.30)	0.12
Rheumatoid arthritis	0.31 (0.14, 0.71)	0.90 (0.41, 2.00)	0.32 (0.14, 0.72)	0.89