Exploring the potential confounder of nitrogen fertilizers in the relationship between pesticide exposures and risk of leukemia: a Poisson regression with two-way fixed-effects analysis

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Dear Editor,

Research discussing potential environmental toxins that may be related to the etiology of childhood leukemia has been growing. The suspected environmental contaminants include solvents, air pollutants, pesticides, and tobacco smoke. Exposure to various pesticides has come under particular scrutiny, with positive associations with childhood leukemia [1]. Poynter et al. [2] conducted a population-based study assessing the association between self-reported chemical exposures and odds of acute myeloid leukemia (AML) and myelodysplastic syndromes (MDS). The authors found no clear association between pesticides and AML or MDS; however, they did report significant associations of AML and MDS with other chemicals, including benzene, vinyl chloride, and fertilizers [2]. While it is certainly important to identify exposure to pesticides as a potential etiological factor in leukemia onset, it is also important to address confounding variables, such as fertilizers, that may be both strongly associated with pesticide use and empirically associated with leukemic outcomes.

We have previously identified environmental exposure to nitrous oxide (N₂O), an agricultural and combustion pollutant, as a likely effect modifier to the proposed relationship between the use of the herbicide, glyphosate, and neurodevelopmental outcomes like attention-deficit hyperactivity disorder (ADHD) [3]. We found that use of glyphosate was closely tied to the use of nitrogen fertilizers in agriculture at a county urbanization level [3]. Therefore, it is possible that pesticide exposures may act as a proxy for air pollutants (i.e., N₂O) directly related to the use of anthropogenic nitrogen in agriculture. Prior studies have identified pre-morbid ADHD and other developmental abnormality in children newly diagnosed with leukemia [4]. Therefore, if environmental N₂O is a trigger for neurodevelopmental disorders, as we have suggested, and developmental abnormalities may precede childhood leukemia, we hypothesize that chronic exposure to N₂O in the environment, and not necessarily pesticide exposures, may foster both neurodevelopmental and hematologic abnormalities.

To investigate the possible association between farm use of nitrogen fertilizers—as the most relevant environmental proxy for N₂O emissions—and hospitalization for blood-related cancers, we have replicated our earlier work [3] using the database from the Healthcare Cost and Utilization Project (HCUP). We conducted a Poisson regression analysis using a two-way fixed-effects model. This approach minimizes the likelihood of omitted variable bias due to unobserved or unmeasured factors that influence the outcome. Briefly, a random variable $Y$ is said to have a Poisson distribution with parameter $\mu$ if it takes integer values $y = 0, 1, 2, \ldots$ with probability.

$$Pr\{Y = y\} = \frac{e^{-\mu} \mu^y}{y!}$$

for $\mu > 0$. The mean and variance of this distribution can be shown to be $E(Y) = \text{var}(Y) = \mu$. We have a sample of $n$ observations of discharges related to blood-related cancers, $y_1, y_2, \ldots, y_n$, which are treated as realizations of independent Poisson random variables, with $Y_{ip} \sim P(\mu_{ij})$.
where $i$ represents a state and $j$ an observation year. We let the logarithm of the mean depend on a vector of time-varying explanatory variables, $x_{ij}$, such that the log-linear model is the following: $\log(\mu_{ij}) = x'_{ij}\beta_1$. We have a multiplicative model for the mean discharges by exponentiation: $\mu_{ij} = \exp(x'_{ij}\beta_1)$. In each case, the exponentiated regression coefficient $\exp(\beta_{1k})$ yields an incidence rate ratio (IRR), which represents a multiplicative effect of the $k$th predictor on the mean. A one log-unit increase in $x_k$ multiplies the mean by a factor $\exp(\beta_{1k})$.

Table 1 shows the results of our Poisson regression analysis using a two-way fixed-effects estimator. The unadjusted model indicates a reduced IRR for hospitalization for leukemia for every one log-unit increase in farm use of nitrogen fertilizers. No other blood cancer (non-Hodgkin's lymphoma or multiple myeloma) was significant in the unadjusted models. When accounting for non-farm use of nitrogen fertilizers as well as the use of pesticides, including atrazine, cypermethrin, dicamba, 2,4-D, glyphosate, and 2-methyl-4-chlorophenoxyacetic acid (MCPA), a one log-unit increase in farm use of nitrogen fertilizers was protective against hospitalization for all three blood cancers. Data for many agricultural states in the USA (such as Iowa, Kansas, and Minnesota) were not available. A third adjusted model excluding the relatively rarely used pesticides, cypermethrin and MCPA, confirms the initial unadjusted model, suggesting that the extreme variability in these two pesticides, in particular, may have contributed to spurious results. However, the reduced risk of hospitalization for leukemia remained statistically significant for all models, lessening the likelihood this result can be attributed to chance. Similar results were noted when using a specific diagnosis of AML in the International Classification of Disease, 9th Revision–Clinical Modification (ICD-9-CM) as the dependent variable.

Including an interaction term of time with farm use of fertilizer heightens the significance of the main effect considerably, while the interaction itself significantly increases annual hospitalization risk for leukemia by 12% (results not shown). These data suggest that a 1-year lagged indicator of farm use of nitrogen fertilizer protects against hospitalization for leukemia, but that compensatory biological mechanisms may be induced over the longer term, soon increasing risk of hospitalizations for leukemia, as supported by Poynter et al. [2].

Biological evidence indicates that $N_2O$ may be protective against leukemic cell growth via the role of $N_2O$ in oxidizing the cobalt ion within cobalamin, inactivating vitamin B12 [5]. Also, $N_2O$ exposure may increase leukocyte DNA damage in patients who underwent surgery, as evidenced by a two-fold increase in the percentage of DNA integrity in the comet tail using digital fluorescence microscopy [6]. The extent of DNA damage was also positively correlated with the duration of $N_2O$ exposure [6]. Therefore, the current epidemiological finding of a significantly protective effect of farm use of nitrogen fertilizers (an environmental proxy of $N_2O$ emissions and exposures) against hospitalization for leukemia is consistent with the prevailing biological evidence. It is interesting to note, however, that elevated blood cobalamin has been acknowledged as a diagnostic marker for leukemia [7], although the significance of this metabolic abnormality in blood cancers is not well characterized. Moreover, the use of methylphenidate, a psychostimulant used to manage premorbid neuropsychiatric conditions like ADHD, has been found to be associated with increased risk of leukemia [8], although evidence of cytogenetic damage

| Outcome | CCSa | Model 1b | Model 2c | Model 3d |
|---------|------|----------|----------|----------|
|         |      | IRR      | 95% CI   | IRR      | 95% CI   | IRR      | 95% CI   |
|         | n = 249 |         |          | n = 90 |         | n = 249 |         |
| Leukemia | 39   | 0.95     | 0.93–0.98 | 0.93  | 0.88–0.98 | 0.95  | 0.93–0.98 |
| NHL     | 38   | 0.98     | 0.94–1.02 | 0.92  | 0.87–0.98 | 0.97  | 0.94–1.01 |
| MM      | 40   | 0.99     | 0.94–1.03 | 0.92  | 0.86–0.99 | 0.98  | 0.94–1.03 |

IRR incidence rate ratio, CI confidence interval

a Using all International Classification of Diseases, Ninth Revision, Clinical Modification (ICD-9-CM) codes included in the Clinical Classification Software (CCS) in HCUP (1997–2007), as described in our prior investigation [3]

b Unadjusted model, farm use of nitrogen fertilizers, as described in our prior investigation [3]

c Controlling for a 1-year lagged, logged measures of non-farm use of nitrogen fertilizers and pesticides, including atrazine, cypermethrin, dicamba, glyphosate, 2,4-D, and 2-methyl-4-chlorophenoxyacetic acid (MCPA), that are associated with arable agriculture crop production (United States Geological Survey, USGS) high estimates, years: 1996–2006 [3]

d Removing cypermethrin and MCPA data due to missingness
attributed to the use of methylphenidate is not consistent [9], suggesting that factors related to the use of psychostimulants and neurodevelopmental impairment, including environmental N₂O exposure, may increase risk of leukemia. These data point to both endogenous and pharmacologic compensatory mechanisms, including leukemic outcomes and methylphenidate use respectively, which may reverse the hematologic and neuropsychiatric effects of chronic N₂O exposure.

The present longitudinal findings provide a more nuanced perspective regarding the significantly increased odds of AML from self-reported exposure to fertilizer [2]. We suggest here that leukemic outcomes, including elevated blood cobalamin levels, may reflect compensatory mechanisms to reverse the hematologic depression from chronic environmental exposure to N₂O emissions associated with nitrogen fertilizer use [10]. Additional investigations are therefore warranted to better characterize the metabolic role of increased serum vitamin B12 in hematologic cancers. As confirmed by Poynter et al. [2], future investigations exploring links between pesticides and leukemia should consider the associated use of nitrogen fertilizers and chronic exposure to related air pollutants, such as N₂O, which has been empirically shown to affect not only leukemic outcomes but also neurodevelopmental abnormality that may often precede a leukemia diagnosis. Therefore, we think that nitrogen fertilizers and their influence on nitrous oxide emissions should be considered.

Abbreviations
ADHD: attention-deficit hyperactivity disorder; AML: acute myeloid leukemia; CCS: Clinical Classification Software; HCUP: Healthcare Cost and Utilization Project; IRR: incidence rate ratio; ICD-9-CM: International Classification of Disease, 9th Revision–Clinical Modification; MDS: myelodysplastic syndromes; MM: multiple myeloma; N₂O: nitrous oxide; NHL: non-Hodgkin’s lymphoma.

Authors’ contributions
KF gathered the data and analyzed the data using appropriate statistical software. Both authors contributed to the writing of the manuscript. Both authors read and approved the final manuscript.

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Competing interests
The authors declare that they have no competing interests.

Availability of data and materials
The datasets generated and/or analyzed during the current study are publicly available in the HealthCare Cost and Utilization Project repository and the United States Geological Survey (persistent weblinks to datasets can be found in the cited prior investigations).

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