

## LETTERS

## Job-related formaldehyde exposure and ALS mortality in the USA

Animal models and in vitro experiments suggest neurotoxic effects of formaldehyde that may be relevant for amyotrophic lateral sclerosis (ALS). Formaldehyde induces neuronal  $\tau$  protein misfolding and aggregation, leading to neuronal apoptosis. Formaldehyde also increases mitochondrial membrane permeability and causes oxidative damage partly by reducing superoxide dismutase activity, mechanisms implicated in ALS.

Studies have had mixed findings regarding formaldehyde exposure and ALS mortality. A large prospective study found an elevated risk that did not quite reach statistical significance, but found a strong dose–response relationship with total years of exposure.<sup>1</sup> Two studies found no significant association,<sup>1 2</sup> although one found a suggestion of elevated risk among the very highly exposed.<sup>2</sup> We examine here the association of ALS mortality with job-related formaldehyde exposure in the National Longitudinal Mortality Study (NLMS), a US-representative cohort with occupation data collected prospectively.

### METHODS

The NLMS is a multistage probability sample of the civilian non-institutionalised population (response rate ~96%). We included the 794 541 men and 674 694 women who were at ages 25+ when surveyed. Participants were asked about their current or most recent job. We used a formaldehyde exposure matrix constructed by industrial hygienists at the National Cancer Institute and previously described.<sup>3</sup> Intensity and probability of formaldehyde exposure were calculated for each occupation and industry, and coded as none, low, medium or high.<sup>3</sup> Intensity reflected the frequency and level of formaldehyde exposure; probability reflected the likelihood of any formaldehyde exposure.

NLMS records were matched to the National Death Index (NDI, 1979–2011) to obtain cause of death. ALS deaths were defined as International Classification of Diseases Ninth and 10th Edition (ICD)-9 335.2 or ICD-10 G12.2 as the underlying or contributing cause. Data were handled and analyses conducted as in prior publications (M G Weisskopf *et al.* Military service and ALS in a population-based cohort; Submitted). Briefly, we estimated HRs using

survival analyses with age as the time meter, adjusted for education, race/ethnicity and income. Participants contributed follow-up time from the time of their survey until time of death or last date of National Death Index (NDI) linkage. We calculated HRs separately for each probability and intensity level, using persons with no exposure as the reference group, separately by sex. We then calculated HRs for each intensity level using the same reference group, and restricting exposed respondents to those with high probability of exposure.

We conducted five sensitivity analyses. As ALS cases are less reliably diagnosed in persons >75 years, we restricted follow-up to age 75. We next conducted four analyses with exclusions that may have improved exposure ascertainment. First, as ALS symptoms may have affected employment, we excluded the first 5 years of follow-up. Second, as the occupation and industry of younger persons is less stable than older persons, we restricted analyses to persons at ages 35–75 at enrolment and ages 50–75 at enrolment. Third, estimated job-related formaldehyde exposure may be less accurate for persons unemployed versus employed at enrolment. We, therefore, restricted the analysis to persons employed at the time of the survey. Additionally, we further adjusted for military service and smoking in the subsample of this data.

### RESULTS

Participants exposed versus unexposed to formaldehyde were slightly poorer, less educated, and less frequently non-Hispanic White (see online supplementary eTable S1). High probability of formaldehyde exposure versus no exposure predicted an almost three times higher rate of ALS mortality in men (table 1). Among women, few had high-exposure-probability jobs and there were no ALS deaths in this category; so the HR was inestimable (see online supplementary eTable S2). Intensity of formaldehyde exposure was less strongly associated with ALS (table 1). High-probability, high-intensity exposure was associated in men with increased rate of ALS mortality (HR=4.43, 95% CI 1.16 to 16.85,  $p<0.05$ ), although there were only two ALS deaths among these highly exposed men. Results were robust to further adjustment for military service and smoking. All men with high-probability, high-intensity exposure were funeral directors. Among men, all sensitivity analyses resulted in higher HR estimates than the main analysis (table 1).

### DISCUSSION

Men in jobs with high probability of exposure versus no formaldehyde exposure had

almost three times greater rate of ALS mortality. We did not find increased risk of ALS in women associated with formaldehyde exposure. Only 99 women in our sample reported jobs with high-probability, high-intensity formaldehyde exposure; thus, our sample of exposed women may have been too small to detect a possible increased risk of ALS. Moreover, all men (N=493) and all but one woman (99%, N=98) in our study in jobs with high-probability high-intensity formaldehyde exposure were funeral directors. In the USA, female versus male funeral directors are more likely to interact with bereaved clients and less likely to perform embalming, where exposure to formaldehyde occurs. Thus, formaldehyde exposure may vary by sex in this profession.

Two prior studies found no association of ALS with job-related formaldehyde exposure. A study of garment workers (geometric mean formaldehyde exposure=0.15 ppm) found no elevated ALS mortality compared with the general population.<sup>4</sup> As garment work does not involve high-probability or high-intensity exposure, these results may not be inconsistent with ours. Funeral directors experience high-intensity and high-probability formaldehyde exposure, with exposure ranging from 0.15 to 9.2 ppm during embalming.<sup>5 6</sup> Additionally, formaldehyde is absorbed through the skin during embalming (at 49.2 mg/h).<sup>6</sup> A second study found no overall association between estimated occupational formaldehyde exposure and ALS. However, in the small subset of participants with the highest exposure to formaldehyde (>60 000 h, N=4 cases, N=4 controls), a large, non-statistically-significant odds for ALS was found (OR=3.0, 95% CI 0.7 to 12.9).

Our results should be interpreted cautiously. Jobs involving both high probability and high intensity of formaldehyde are relatively uncommon in the USA, and ALS is also rare; there were only two ALS deaths among men in such jobs. Moreover, we did not find a dose–response association between formaldehyde exposure and ALS. Formaldehyde exposure was estimated from a single report at enrolment. This single job report likely did not accurately capture lifetime exposure. As non-differential error in exposure classification typically leads to attenuation of the true association of exposure with disease, our estimated HRs could have been attenuated and any trend obscured.

In addition to formaldehyde, funeral directors are exposed to other chemicals used in embalming, as well as to viral, bacterial and prion pathogens. Thus, further study of the association of ALS with high

**Table 1** Adjusted HRs\* and 95% CIs for ALS mortality by level of occupational formaldehyde exposure, National Longitudinal Mortality Study, men ages 25 years and older, 1973–2011

	Respondents	Person-years	ALS deaths	HR (95% CI)*
<b>Intensity</b>				
Unexposed	607 416	9 815 195	372	1.0 (Reference)
Low	97 301	1 641 068	55	0.99 (0.74 to 1.30)
Medium	86 766	1 427 789	43	0.63 (0.44 to 0.90)
High	3058	46 188	2	1.53 (0.40 to 5.80)
<b>Intensity, exposed restricted to probability=high</b>				
Unexposed	607 416	9 815 195	372	1.0 (Reference)
Low	0	0	0	–
Medium	361	6954	0	–
High	493	8539	2	4.43 (1.16 to 16.85)
<b>Probability</b>				
Unexposed	607 416	9 815 195	372	1.0 (Reference)
Low	98 228	1 681 862	51	0.85 (0.63 to 1.15)
Medium	88 043	1 417 659	47	0.76 (0.54 to 1.06)
High	854	15 493	2	2.98 (0.78 to 11.30)
<b>Probability, excluding first 5 years of follow-up</b>				
Unexposed	492 489	9 501 370	331	1.0 (Reference)
Low	82 155	1 637 260	47	0.87 (0.64 to 1.18)
Medium	71 598	1 372 603	39	0.77 (0.54 to 1.10)
High	758	15 230	2	3.49 (0.92 to 13.26)
<b>Probability, follow-up to age 75 only</b>				
Unexposed	604 116	9 794 521	332	1.0 (Reference)
Low	97 959	1 680 822	41	0.79 (0.57 to 1.11)
Medium	86 760	1 407 671	40	0.66 (0.44 to 0.99)
High	832	15 315	2	4.13 (1.09 to 15.69)
<b>Probability, respondents 35≤age≤75 at enrolment</b>				
Unexposed	427 530	6 493 151	332	1.0 (Reference)
Low	65 736	1 046 575	47	0.91 (0.67 to 1.24)
Medium	60 675	921 096	43	0.79 (0.56 to 1.12)
High	568	9731	2	3.41 (0.89 to 13.01)
<b>Probability, respondents 50≤age≤75 at enrolment</b>				
Unexposed	175 376	2 441 370	197	1.0 (Reference)
Low	25 611	377 534	31	1.00 (0.67 to 1.49)
Medium	27 247	393 151	27	0.75 (0.47 to 1.19)
High	270	4125	2	4.76 (1.16 to 19.49)
<b>Probability, respondents employed at enrolment</b>				
Unexposed	548 645	8 876 712	331	1.0 (Reference)
Low	85 041	1 459 452	45	0.88 (0.65 to 1.21)
Medium	76 249	1 238 082	38	0.71 (0.50 to 1.03)
High	770	14 063	2	3.26 (0.86 to 12.38)

\*Adjusted for race/ethnicity, education and household income as a percentage of the poverty line.  
ALS, amyotrophic lateral sclerosis.

questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. NJJ had full access to the data and takes responsibility for accuracy of the data analyses.

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## REFERENCES

- Weisskopf MG, Morozova N, O'Reilly EJ, et al. Prospective study of chemical exposures and amyotrophic lateral sclerosis. *J Neurol Neurosurg Psychiatry* 2009;**80**:558–61.
- Fang F, Quinlan P, Ye W, et al. Workplace exposures and the risk of amyotrophic lateral sclerosis. *Environ Health Perspect* 2009;**117**:1387–92.
- Wang R, Zhang Y, Lan Q, et al. Occupational exposure to solvents and risk of non-Hodgkin lymphoma in Connecticut women. *Am J Epidemiol* 2009;**169**:176–85.
- Pinkerton LE, Hein MJ, Meyers A, et al. Assessment of ALS mortality in a cohort of formaldehyde-exposed

levels of formaldehyde exposure and among funeral directors is warranted.

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- garment workers. *Amyotroph Lateral Scler Frontotemporal Degener* 2013;14:353–5.
- 5 Williams TM, Levine RJ, Blunden PB. Exposure of embalmers to formaldehyde and other chemicals. *Am Ind Hyg Assoc J* 1984;45:172–6.
  - 6 Beoniger MF, Stewart P. Biological markers for formaldehyde exposure in mortician students: extent of exposure. In: National Institute for Occupational Safety and Health, ed. Cincinnati, Ohio: US Department of Health and Human Services, 1992:45.