

Exploratory breath analyses for assessing toxic dermal exposures of firefighters during suppression of structural burns

Joachim D Pleil¹, Matthew A Stiegel² and Kenneth W Fent³

¹ Human Exposure and Atmospheric Science Division, NERL/ORD, US Environmental Protection Agency, Research Triangle Park, NC, USA

² ORISE/EPA Research Fellow, Gillings School of Global Public Health, University of North Carolina, Chapel Hill, NC, USA

³ Division of Surveillance, Hazard Evaluations, and Field Studies, National Institute for Occupational Safety and Health (NIOSH), Cincinnati, OH, USA

E-mail: pleil.joachim@epa.gov

Received 11 March 2014, revised 27 May 2014

Accepted for publication 5 June 2014

Published 4 September 2014

Abstract

Firefighters wear fireproof clothing and self-contained breathing apparatus (SCBA) during rescue and fire suppression activities to protect against acute effects from heat and toxic chemicals. Fire services are also concerned about long-term health outcomes from chemical exposures over a working lifetime, in particular about low-level exposures that might serve as initiating events for adverse outcome pathways (AOP) leading to cancer. As part of a larger US National Institute for Occupational Safety and Health (NIOSH) study of dermal exposure protection from safety gear used by the City of Chicago firefighters, we collected pre- and post-fire fighting breath samples and analyzed for single-ring and polycyclic aromatic hydrocarbons as bioindicators of occupational exposure to gas-phase toxicants. Under the assumption that SCBA protects completely against inhalation exposures, any changes in the exhaled profile of combustion products were attributed to dermal exposures from gas and particle penetration through the protective clothing. Two separate rounds of firefighting activity were performed each with 15 firefighters per round. Exhaled breath samples were collected onto adsorbent tubes and analyzed with gas-chromatography-mass spectrometry (GC-MS) with a targeted approach using selective ion monitoring. We found that single ring aromatics and some PAHs were statistically elevated in post-firefighting samples of some individuals, suggesting that fire protective gear may allow for dermal exposures to airborne contaminants. However, in comparison to a previous occupational study of Air Force maintenance personnel where similar compounds were measured, these exposures are much lower suggesting that firefighters' gear is very effective. This study suggests that exhaled breath sampling and analysis for specific targeted compounds is a suitable method for assessing systemic dermal exposure in a simple and non-invasive manner.

Keywords: adverse outcome pathway, exposome, firefighter, JP-8 jet fuel, PAHs, benzene, heatmap

(Some figures may appear in colour only in the online journal)

1. Introduction

Firefighters wear specialized gear designed primarily to protect against the immediate dangers of fire associated with heat, asphyxiation, and acute effects from toxic vapors, gases, and

particles from combustion sources. This article addresses a secondary concern, which is the potential for lower level toxic exposures that may create longer-term adverse health outcomes. Such exposures pose a health risk by initiating adverse outcome pathways (AOPs) at the molecular or cellular level

and are experienced repeatedly by the community of 330 000 career fire fighters and 770 000 volunteer fire fighters in the United States (NFPA 2011). As such, it is important to gain detailed exposure data from representative single events to predict the overall risk to the human systems biology for long-term latency disease from repeated short-term exposures (Pleil 2009, Pleil and Sheldon 2011). Furthermore, the availability of *in vivo* biomarker specimens is valuable in supplementing the overall exposure assessment framework and the toxicity data derived from *in vitro* (molecular and cellular level) and *in silico* (computational models) research of suspected human carcinogens (Pleil *et al* 2012, Sobus *et al* 2011, Tan *et al* 2012).

The US National Institute for Occupational Safety and Health (NIOSH) conducted a study of dermal exposures to and biological uptake of carcinogenic compounds in fire-fighters during fire suppression. Firefighters in this study wore the highest level of dermal and respiratory protection currently available to them. This study included ambient measurements, urinary biomarkers, dermal swabs, and assessments of protective fire-fighting turnout gear (Fent *et al* 2013). As part of this study, US Environmental Protection Agency (EPA) researchers collaborated with NIOSH in assessing exhaled breath biomarkers of fire exposure represented by aromatic and polycyclic aromatic hydrocarbons (PAHs). Breath analysis has become a useful procedure for a variety of environmental and clinical health investigations (Davis *et al* 2013, Dweik RA 2011, Miekisch *et al* 2012, Pleil 2008) and has been featured in recent conferences and review articles as a mainstream technology (Amann 2009, Beauchamp and Pleil 2013, Braun, Gmachl and Dwaek 2012, Corradi and Mutti 2012, Pleil and Hanzel 2012, Pleil *et al* 2013, Risby and Solga 2006). Furthermore, breath has joined blood and urine as a primary human biofluid in decoding the human exposome (Pleil *et al* 2011, Sinues *et al* 2013).

The purpose of the breath portion of the study was to provide objective biomarker evidence of short-term exposures and was conducted as a 'targeted' rather than a 'discovery' assessment (Pleil and Stiegel 2013). That is, the analytes were chosen *a priori* to optimize sensitivity and specificity. As such, we restricted ourselves to certain compounds that are presumed indicative of the totality of exposure to products of incomplete combustion; this approach has been used for environmentally derived samples for assessments of other fire related exposures including from the 9/11 New York City World Trade Center (WTC) disaster (Lorber *et al* 2007, Pleil *et al* 2004, Pleil *et al* 2006). We note that there are previous breath based studies of firefighters for assessing carbon monoxide (CO) exposures (Cone *et al* 2005, Dunn *et al* 2009, Miranda *et al* 2012), but that this is the first study wherein organics exposures were directly assessed using breath biomarkers.

This article describes the application and results from breath analyses of fire fighters engaged in scripted (planned exposure) activities. Breath samples were collected before, shortly after, and 6 h after specific firefighting tasks in a realistic training environment. The breath results indicated that some amount of biological uptake of toxic materials occurred despite the use of dermal and respiratory protection. We compared these results to other breath studies of jet fuel exposures

to gain context of occupational exposures to similar compounds. Exhaled breath analysis was found to be a valuable and noninvasive tool for assessing exposures of firefighters to toxic chemicals derived from incomplete combustion.

2. Methods

2.1. Study design

The study was comprised of two rounds of controlled structure burns 1 year apart; each round consisted of 3 days of scripted firefighting activity with five firefighters studied per day. As such, there were 30 distinct subject-days of data gathered; twelve of the firefighters from round 1 repeated as subjects in round 2. Each day of firefighting activity consisted of a scripted test burn that took place in the morning with segments of fire entry, knockdown, and overhaul. Firefighting activity varied in time from 15 to 30 min depending on the heat and intensity of each of the fires. The firefighters wore protective ensembles for structural firefighting that were compliant with the National Fire Protection Association 1971/1981 standards (NFPA 2007a, 2007b) and did not remove their ensembles until the controlled burns had been completely extinguished and they were 30 meters or more away from the burn structure in a clean air environment. We note that any residual off-gassing of the gear that may have mixed with the ambient air during gear removal could have constituted a brief (minute-long) trace-level inhalation exposure. However, post-work exhaled breath monitoring was initiated after about 10 min delay to assure that measurements reflected only longer-term systemic exposures from the fires. This is reasonable as uptake and elimination kinetics are symmetric and the effects of a brief (few breaths) exposure are eliminated in the same time frame (Kim *et al* 2007, Pleil and Lindstrom 1998). The controlled burns were conducted by the Illinois Fire Service Institute at the University of Illinois—Urbana-Champaign. The subset of samples described herein were comprised of end-tidal breath and served to confirm specific exposures. A detailed description of each round-day is available in the NIOSH study report (Fent *et al* 2013).

2.2. Breath sampling and analysis

Breath samples were collected from each firefighter approximately 1 h before the activity, about 10 min after the conclusion of the activity, and then 6 h post activity. Subjects forcefully exhaled an entire breath through a Markes International Bio-VOC™ sampler wherein the last 127 ml of end-tidal air were retained, and subsequently transferred onto a Markes Carbograph 2TD/Carbograph 1TD thermal desorption tube (Markes International, Ltd, Llantrisant, UK). The tubes were shipped to the US EPA laboratories at Research Triangle Park, NC, USA for analysis. Tubes were sealed and stored under refrigeration from sampling, through shipment, and until immediately before analysis according to our standard internal procedures.

Adsorbent tubes were analyzed via methods developed at EPA for a variety of VOCs in all types of gas-phase matrices and sampling containers (Hubbard *et al* 2009, Pleil 2008,

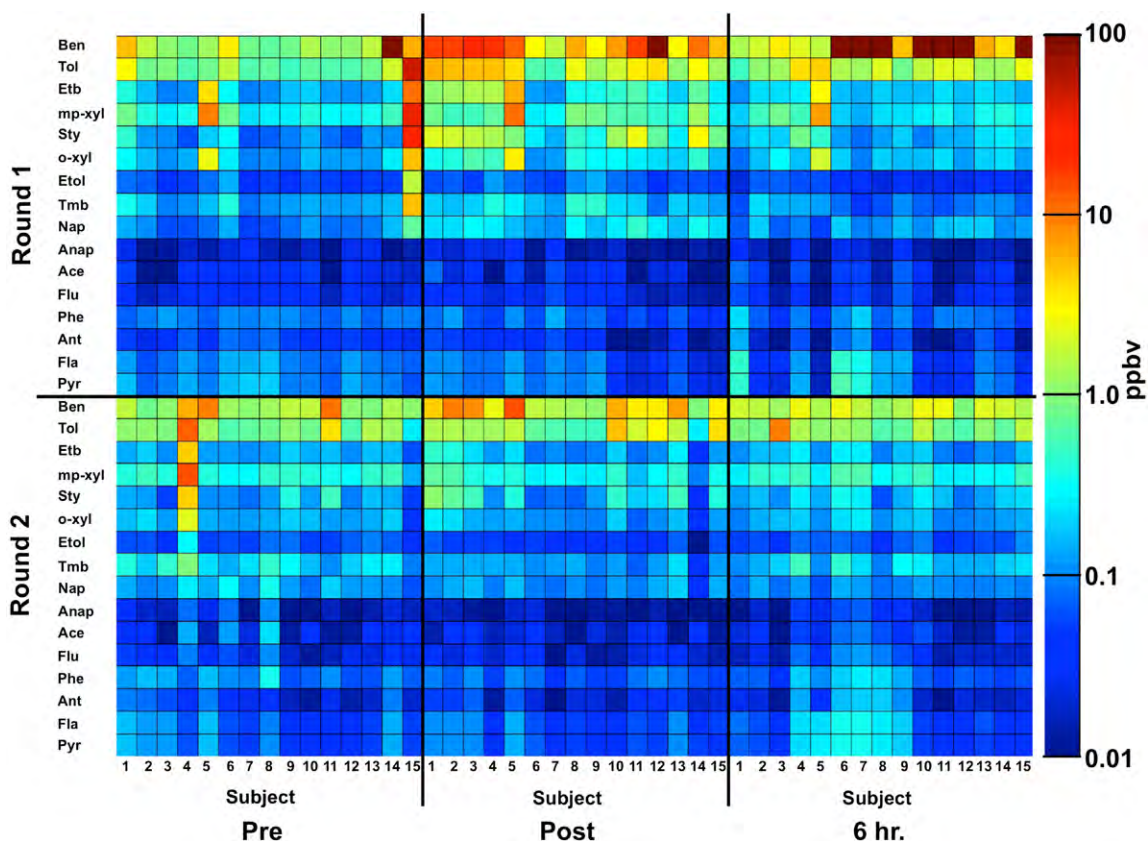


Figure 1. Heatmap of exhaled breath concentrations for NIOSH firefighters dermal exposure study grouped by exposure period (*x*-axis) and by experimental round and compounds (*y*-axis). Patterns indicate no major changes.

Pleil and Lindstrom 1995). Specifically, for this project we used an Ultra TD autosampler coupled with a Unity thermal desorber for the adsorbent tubes (Markes International, Ltd, Llantrisant, UK) coupled to a GC/MS system (6890N GC, 5973N MS, Agilent Technologies, Palo Alto, CA). The samples were analyzed in selected ion monitoring (SIM) mode with optimized ion groups and dwell times to achieve a sampling rate of approximately 2 Hz and a minimum of 10 points per GC peak (Pleil *et al* 1991). To insure that the peaks of interest were correctly identified, the peak area for each analyte was calculated and compared for both primary and secondary ions. Instrumentation was externally calibrated for all targeted analytes using certified standards (Accustandard, New Haven, CT, USA). Standard instrument and tube blank analyses were performed to set the zero and span calibration slopes for quantitation. The levels of quantitation (LoQ) for the aromatic hydrocarbons are <0.09 ppbv, and the LoQs for the 2 to 4-ring PAHs are in the range of 0.004–0.01 ppbv.

2.3. Human subjects

Firefighter subjects were recruited as volunteers from the Chicago Fire Department. Inclusion criteria were non-smoking males, 45 years of age or younger, with instructor level firefighter qualifications. Subjects were instructed to refrain from eating charbroiled food and avoid second hand tobacco smoke for at least 2 days prior to their participation. Their respective fire chiefs ensured that volunteers had at

least 1 day off before reporting to the study. Subjects participated with informed consent under Institutional Review Protocol HSRB 10-DSHEFS-03 administered by NIOSH.

2.4. Mathematical analysis

This set of experiments has a complex nested data structure: 2 rounds, 3 days per round, 5 subjects per day, 3 samples per subject. We chose to composite the days within rounds into individual data sets (15 subjects each) to boost statistical power. Because twelve of the subjects appeared in both rounds, we had the opportunity to treat their results as repeat measures to gain insight into within-subject and between-subject variance components (Pleil 2009, Pleil and Sobus 2013, Sobus *et al* 2010). In addition to summary statistics comparing the temporal profiles of exhaled organics (i.e. pre, post and 6 h post), we also assessed the individual subjects profiles by normalizing the post-work samples by the pre-work and 6 h post-work samples as a baseline control. Data were compiled and analyzed using MS Excel (Microsoft Corp., Redmond, CA); graphics were created using GraphPad Prism (GraphPad Software Inc., La Jolla, CA).

3. Results

3.1. General observations

Figure 1 presents all of the firefighter breath data in a heatmap. The data are arranged horizontally along the *x*-axis

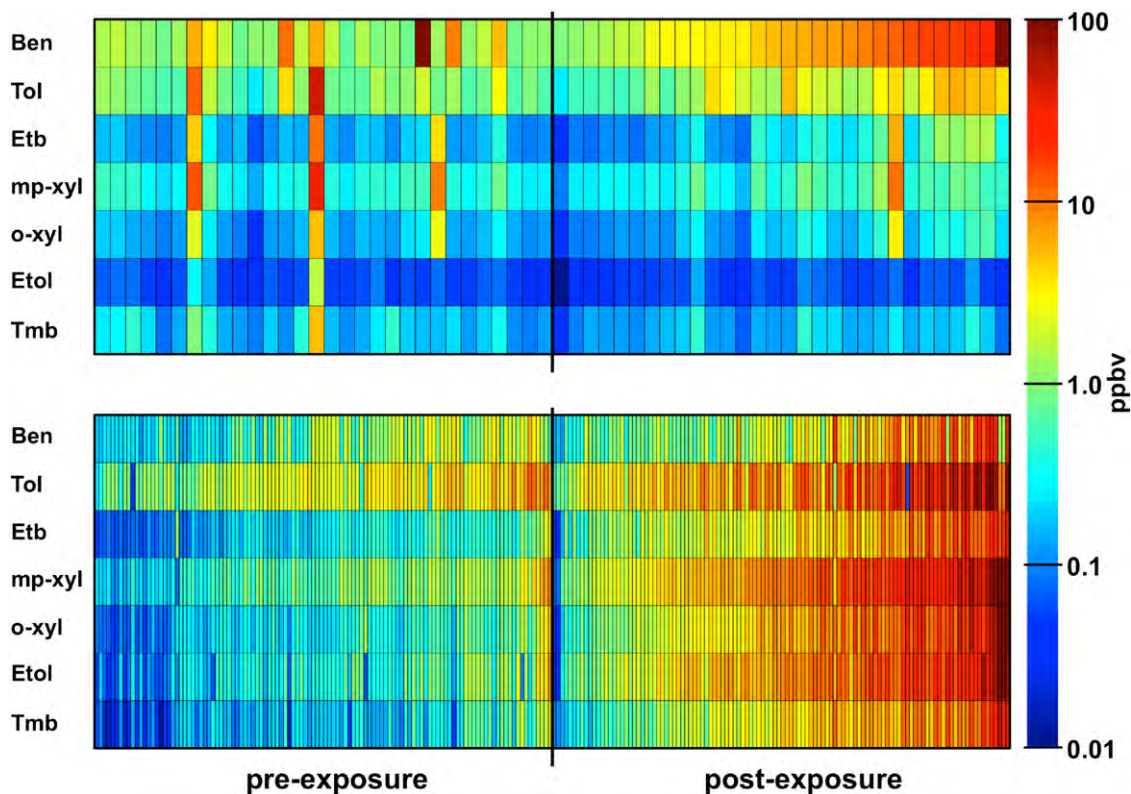


Figure 2. Direct heat map comparisons of occupational exposures to gas-phase hydrocarbons (single-ring aromatic compounds) as measured in exhaled breath. Upper section shows the heat map of fire fighters’ exposures demonstrating a modest increase in benzene, and toluene immediately after the exposure. Lower section shows the corresponding heat map for Air Force subjects’ exposures demonstrating across the board increases in exhaled aromatic hydrocarbons.

as three blocks (by subject number) representing pre-, post and 6 h post measurements. The vertical axis is arranged as two blocks representing rounds 1 and 2 with internal structure of compounds arranged in order from top to bottom by elution on the GC column. The corresponding fields in the *x-y* plane are color coded to represent the concentration of analytes in the breath according to the color scale (red = high to blue = low) found at the far right column of the graphic. From this observational tool, we see that there are large variances among subjects, there are no obvious major trends between time points, and most of the apparent outliers occurred during round 1. Furthermore, the less volatile compounds (3-ring PAHs) have much lower levels in the breath than the more volatile single ring aromatics. The outlier group in the round 1, 6 h post-work benzene levels (upper right side corner) comprised of subjects 6, 7, 8, 10, 11, 12, and 15 are most likely due to some external exposures that were unnoticed by the field operatives; this was not observed in the second round. We believe that these exposures were unrelated to firefighting, as the pattern of the other compounds seems to fit in well with the remaining data. We also note the occasional high outlier series of the volatiles e.g. round 1, pre-work subject 15, round 2, pre-work subject 4 and round 1, 6 h post-work subject 5. We note that the data appear log-normally distributed within compound, as expected, and as observed in the log nature of the color code. We confirmed this with subsequent calculations described below.

For context, we extracted results from previously published studies of jet fuel exposures in the US Air Force (USAF) (Pleil *et al* 2011, Pleil, Smith and Zelnick 2000). The USAF studies measured volatile JP-8 jet fuel constituents that included n-alkanes and single-ring aromatic compounds; for the comparison, we chose only those compounds that overlapped with the present study: benzene (Ben), toluene (Tol), ethylbenzene (Etb), m,p-xylene (mp-xyl), o-xylene (o-xyl), ethyl toluene (Etol) and 1,3,5-trimethylbenzene (Tmb). Figure 2 shows the direct comparisons in a two-part heatmap: the upper portion shows the pre- and post-exposure comparison for the firefighters, and the lower portion shows the pre- and post-exposure comparison for the USAF personnel. The heat map comparison shows very distinctly that the occupational exposures in the USAF studies were much more pronounced. In both the firefighter and Air Force studies the subjects wore supplied air respiratory protection; however, due to the confined spaces within the aircraft and concerns for explosion risk, the Air Force workers wore cotton overalls, and so, were much more likely to have dermal exposures from vapors and accidental contact with fuel. Figures 3(a) and (b) show typical exposure scenarios for both studies and documents the personal protective differences.

Figures 3(a) and (b) (below) show photographs of representative exposure scenarios for both studies and documents the personal protective gear. We see that the firefighters’ turnout gear is much more substantive than the aircraft



Figure 3. Contrasting exposures and general surroundings from firefighter (a) and aircraft mechanics (b) studies. Visually, fire exposures appear more widespread and intense and firefighters’ gear more heavily protective. Both occupations employ forced air respirators (albeit with different types); the difference may be that firefighters’ gear is more effective against dermal exposures.

mechanics’ gear (especially against dermal exposures), presumably because their workspace and general surroundings are much more hazardous and contaminated. These pictures are intended only to give the reader a comparative view of the occupational scenarios; due to the highly variable exposures in both cases, we cannot provide direct ambient air comparisons. However, we (NIOSH) made a series of total PAHs measurements during the structural burns and found a median value of about 375 ppmv with an overall range of about 10 ppmv to 3000 ppmv. We note that the Air Force environments were considered to be below levels of acute effects such as asphyxiation, inflammation, or neurological damage

without protective gear, unlike the atmospheres encountered in structural fire.

3.2. Statistical analysis: overall trends

We calculated summary statistics for rounds 1 and 2 of all raw data for each of the 15 compounds and each of the three time points. As seen in the heat map in figure 1, there are a few inexplicable outliers in the measurements that might skew the mathematical results of the raw data; we subsequently removed these outliers from the data set if they exceeded the 99th percentile based on geometric mean (GM) and geometric

Table 1. Summary statistics for exhaled breath concentrations (ppbv).

Concentration: ppbv	Pre-exposure				Post-exposure				6 h post-exposure			
	Compound Round 1	<i>n</i>	GM	GSD	95%	<i>n</i>	GM	GSD	95%	<i>n</i>	GM	GSD
benzene	14	1.469	0.620	5.262	14	7.713	0.722	36.551	9	4.152	1.090	42.027
toluene	14	0.869	0.449	2.330	15	2.323	0.580	9.975	15	1.580	0.471	4.622
ethylbenzene	13	0.152	0.366	0.364	14	0.447	0.528	2.095	14	0.180	0.292	0.284
m,p-xylene	13	0.355	0.344	0.759	14	0.554	0.354	1.245	14	0.287	0.367	0.689
styrene	14	0.114	0.420	0.345	15	0.826	0.593	4.728	14	0.167	0.357	0.376
o-xylene	13	0.149	0.329	0.291	14	0.278	0.382	0.717	14	0.163	0.323	0.311
4-ethyltoluene	14	0.055	0.306	0.119	15	0.070	0.282	0.131	15	0.037	0.312	0.084
1,3,5-trimethylbenzene	14	0.148	0.313	0.333	15	0.203	0.323	0.483	15	0.090	0.310	0.198
naphthalene	14	0.088	0.269	0.168	15	0.217	0.272	0.426	15	0.117	0.280	0.242
acenaphthalene	15	0.016	0.247	0.036	15	0.014	0.303	0.047	15	0.014	0.336	0.056
acenaphthene	15	0.024	0.330	0.095	15	0.022	0.353	0.096	15	0.023	0.434	0.150
fluorene	15	0.029	0.203	0.054	15	0.026	0.238	0.063	15	0.028	0.274	0.090
phenanthrene	15	0.089	0.164	0.126	15	0.065	0.205	0.138	15	0.079	0.233	0.213
anthracene	15	0.046	0.200	0.094	15	0.024	0.322	0.118	15	0.029	0.403	0.213
fluoranthene	15	0.099	0.160	0.170	15	0.058	0.236	0.202	15	0.078	0.361	0.596
pyrene	15	0.110	0.170	0.209	15	0.063	0.239	0.225	15	0.081	0.385	0.693
Compound Round 2	<i>n</i>	GM	GSD	95%	<i>n</i>	GM	GSD	95%	<i>n</i>	GM	GSD	95%
benzene	15	1.779	0.708	8.131	15	3.280	0.733	16.002	14	1.696	0.427	3.064
toluene	14	0.954	0.482	2.912	15	1.264	0.596	5.708	13	0.966	0.376	1.868
ethylbenzene	14	0.133	0.308	0.230	15	0.136	0.445	0.464	14	0.143	0.352	0.318
m,p-xylene	14	0.365	0.324	0.696	15	0.318	0.375	0.792	14	0.380	0.321	0.715
styrene	14	0.126	0.437	0.411	15	0.215	0.613	1.305	14	0.159	0.379	0.400
o-xylene	14	0.138	0.327	0.268	15	0.123	0.358	0.281	14	0.143	0.319	0.265
4-ethyltoluene	15	0.066	0.318	0.154	15	0.045	0.307	0.098	14	0.064	0.274	0.112
1,3,5-trimethylbenzene	15	0.244	0.363	0.722	15	0.120	0.290	0.235	14	0.192	0.336	0.493
naphthalene	15	0.088	0.269	0.168	15	0.112	0.277	0.226	14	0.099	0.243	0.159
acenaphthalene	13	0.017	0.290	0.051	15	0.011	0.292	0.033	14	0.021	0.363	0.094
acenaphthene	14	0.028	0.368	0.135	15	0.020	0.248	0.048	14	0.032	0.324	0.123
fluorene	14	0.033	0.257	0.092	15	0.022	0.226	0.048	14	0.035	0.321	0.151
phenanthrene	14	0.095	0.195	0.185	15	0.069	0.188	0.125	14	0.097	0.247	0.292
anthracene	15	0.035	0.285	0.137	15	0.028	0.248	0.085	14	0.042	0.409	0.317
fluoranthene	15	0.072	0.208	0.198	15	0.063	0.190	0.148	14	0.097	0.286	0.483
pyrene	15	0.072	0.213	0.207	15	0.063	0.196	0.157	14	0.097	0.308	0.553

standard deviation (GSD) calculations and recalculated the summary statistics. The summary statistics (*n*, GM, GSD, 95% value) are presented by rounds 1 and 2 and the three time points: pre-, post-, and 6 h post-exposure. As we saw in the qualitative patterns in the figure 1 heatmap, the summary statistics confirm that there are no obvious overall trends. We present these results primarily to show the concentrations expected in such a scenario, and what could be considered ‘unremarkable’ based on the pre- and 6 h post-exposure levels. We further note that all measurements were above the LoQs for their respective compounds.

We subsequently reduced the composite data from table 1 to assess statistical significance of any of the time point differences within the overall scatter of the results. Table 2 shows one-sided significance ($\alpha = 0.05$) for an increase of the post-exposure value over the pre-exposure value (post > pre) and the post exposure value over the 6 h post-exposure value (post > 6h). The assumption that 6 h are sufficient to eliminate exposures from the 1 h firefighting activity is reasonable as breath uptake and elimination kinetics are generally symmetric (Pleil and Lindstrom 1998). Research has also shown that dermal exposures have similar kinetics for non-polar compounds and naphthalene has been used as the marker for fuels research (Kim *et al* 2007). Results are presented for

each round separately and as a compilation of both rounds. The measurements were treated as independent, so we did not administer multiple test corrections.

We see that overall there is a trend that most of the single-ring aromatics and naphthalene have a statistically significant exposure effect in at least one category. The 3- and 4-ring PAHs do not show any significant trends; this is expected, as short-term exposures are generally not sufficient to overcome the general background of systemic PAHs accumulated from other environmental sources, especially food. Previous studies have shown that short term (2h) scripted exposures to diesel exhaust e.g. do not statistically increase the higher PAHs, and that overall, blood and plasma samples in randomly selected healthy subjects have similar ranges of onboard PAHs (Ghio *et al* 2012, Sobus *et al* 2010). Detailed analyses with human metadata and multiple studies have confirmed this observation (Lu *et al* 2014).

3.3. Statistical analysis: individual (subject) trends

We considered that the overall variability among subjects might have obscured individual (auto-correlated) changes. We explored this phenomenon by normalizing each post exposure measurement per individual by the average of their own

Table 2. Exposure significance by compound and experimental round ($\alpha = 0.05$).

Compound	Round 1		Round 2		Aggregate	
	post > pre	post > 6h	post > pre	post > 6h	post > pre	post > 6h
benzene	yes	—	yes	yes	yes	yes
toluene	yes	—	—	—	yes	—
ethylbenzene	yes	yes	—	—	yes	yes
m,p-xylene	yes	yes	—	—	—	yes
styrene	yes	yes	yes	—	yes	yes
o-xylene	yes	yes	—	—	—	—
4-ethyltoluene	yes	yes	—	—	—	—
1,3,5-trimethylbenzene	yes	yes	—	—	yes	—
naphthalene	yes	yes	yes	—	yes	yes
acenaphthalene	—	—	—	—	—	—
acenaphthene	—	—	—	—	—	—
fluorene	—	—	—	—	—	—
phenanthrene	—	—	—	—	—	—
anthracene	—	—	—	—	—	—
fluoranthene	—	—	—	—	—	—
pyrene	—	—	—	—	—	—

Table 3. Individual firefighters, by round, exhibiting greater than 50% increase in exhaled breath concentration over their own baseline values.

Compounds	Round 1	Round 2
benzene	2, 4, 8, 11, 12, 13, 15	3, 4, 5, 7, 12, 13, 14
toluene	2, 7, 8, 9, 11, 13	3, 4, 14
ethylbenzene	2, 5, 7, 8, 9, 11, 13, 14	1, 5, 13, 14
m,p-xylene	2, 5, 6, 7, 11, 13	1, 14
styrene	1, 2, 3, 4, 5, 6, 7, 8, 9, 11, 13, 14	1, 3, 4, 5, 7, 12, 13, 14
o-xylene	2, 5, 7, 8, 11, 13, 14	1, 14
4-ethyltoluene	2, 4, 5, 6, 12	1
1,3,5-trimethylbenzene	2, 4, 5, 6, 7, 12	1
naphthalene	2, 3, 4, 6, 7, 8, 11, 12, 13, 14	5, 7
acenaphthalene	4, 8, 9, 10	6
acenaphthene	4, 9, 10	2, 5
fluorene	4, 9, 10	5
phenanthrene	8, 10	5
anthracene	4, 8	5
fluoranthene	4, 8	5
pyrene	4, 8	5

pre-exposure and 6h post-exposure values (under the assumption that 6h is sufficient to wash out the immediate increments from the fire fighting activity). We used the data (table 1) from which the extreme outliers had already been removed to harmonize the comparisons. We found that some firefighters had significant increases in breath biomarkers measured immediately post-work whereas others had no statistical changes. Table 3 shows the results at the individual level. For this exercise, we arbitrarily considered the individual’s exposure to be of interest if the post-value is at least 50% higher than the established individual baseline.

Here we see that certain firefighters were more consistently exposed than others. We also see that round 2 had appreciably fewer firefighters with relevant exposures. Overall, firefighters #’s 2, 4, 8, 11, and 13 in round 1, and #’s 1, 5, and 14 in round 2 exhibited significant exposure hits for 6 or more compounds, whereas #’s 1 and 3 in round 1, and #’s 2, 6, 7, 8, 9, 10, and 15 in round 2 had insignificant incremental exposures (1 or none exceedances). We note that firefighters numbered 1 through 12 are the same individuals in both rounds; these data indicate that exposures seem to be driven by the particular events, rather than by individual traits or behaviors. We also suggest that the specific training exercises, even within

the same experimental round, are highly variable as fires (even under controlled burn conditions) are subject to many different factors. These observations are more robustly evaluated in the next section.

3.4. Statistical analysis: intra-class correlations

An important feature of environmental and occupational exposure science is the assessment of within- and between-subject variance components (σ_w^2 and σ_b^2 , respectively). This helps discern how to mitigate exposure exceedances and to get an estimate of average exposures that influence long-term risk. We have discussed this in detail elsewhere (Pleil and Sobus 2013a, 2013b), but briefly, when repeat measures are available, one can estimate the intra-class correlation co-efficient (ICC), which is defined as $ICC = \sigma_b^2 / (\sigma_w^2 + \sigma_b^2)$ and takes on a value between 0 and 1. When ICC is near 1, most of the variability is between subjects and implies that repeated measures for any individual yield approximately the same value; when ICC is closer to 0, repeated measurements will be fairly random regardless of the individual measured.

In this study, 12 of the 15 total subjects were common to rounds 1 and 2, and so we could treat them as having

Table 4. ICC calculations using round 1 and round 2 as repeat measures.

Compounds	pre	post	6h
benzene	0.000	0.811	0.840
toluene	0.030	0.505	0.175
ethylbenzene	0.000	0.899	0.674
m,p-xylene	0.000	0.789	0.000
styrene	0.011	0.876	0.168
o-xylene	0.000	0.881	0.620
4-ethyltoluene	0.252	0.789	0.733
1,3,5-trimethylbenzene	0.732	0.822	0.795
naphthalene	0.792	0.890	0.229
acenaphthalene	0.000	0.563	0.000
acenaphthene	0.000	0.331	0.000
fluorene	0.000	0.649	0.000
phenanthrene	0.000	0.000	0.000
anthracene	0.571	0.000	0.000
fluoranthene	0.731	0.134	0.000
pyrene	0.816	0.344	0.000

independent ‘repeat measures’ 1 year apart. We arranged the data to pair round 1 to round 2 measurements for all compounds by sampling time point.

From table 4, we interpret low ICC ($ICC < 0.4$) as indicating relatively weak year-to-year correlation, and high ICC ($ICC > 0.5$) as indicating stronger year-to-year correlation within the respective sampling time-points. These results are based on non-dimensional analysis (performed in log-space), and so absolute exposure levels are divided out; what we can interpret is the relative influences from environmental and occupational (firefighting) sources. We see that the post-time point ICC values for the single-ring aromatics (benzene through 1,3,5-trimethylbenzene) and some of the PAHs (naphthalene, acenaphthalene and fluorine) are > 0.5 , indicating that these compounds affected firefighters similarly during rounds 1 and 2 exposures. The corresponding low ICC values in the pre column indicate that the environmental exposures are fairly random for these subjects. This is confirmed by the nature of exposure to volatiles (single-ring aromatics) wherein the elimination half-lives are relatively short. What is of particular interest is the handful of high ICC values in the 6h column; we would have expected these values to revert to their respective pre levels. These unexpected results indicate that there were likely some ongoing residual exposures of benzene, ethylbenzene, etc from unknown sources that were too subtle to be seen in the qualitative heatmaps in figure 1.

4. Conclusions

This study demonstrated that exhaled breath biomarkers represent a useful technology for assessing occupational exposures of firefighters to products of combustion, especially of toxic aromatic compounds. Of particular note is the novel application to the less volatile 3- and 4-ring PAHs (ranging from acenaphthalene to pyrene) that generally require blood or urine sampling. We found that background (pre-exposure and 6h post-exposure) levels are within the sensitivity of the methodology, that the single-ring compounds have overall higher levels in breath than the PAHs,

and that the breath borne concentrations can range from low parts per trillion (pptv) to 100’s of parts per billion (ppbv) (figure 1, table 1).

We found that there are immediate post-exposure exceedances above an expected baseline of many of the gas-phase and semi-volatile aromatic compounds (table 2). Furthermore, we found that individual firefighters have widely varying exposure profiles (figure 1, table 3) and we attribute this to specific situations, activities, timing and microenvironments from the inherent heterogeneity of fires. From the calculations of ICC’s using round 1 and round 2 as repeat measures, we conclude that the exposures are most likely driven by situational (between-subject) rather than within-subject variance components, and that there were some residual exposures from unknown sources occurring during or just prior to the 6-h post-work period.

In figure 2 we demonstrate that the protective gear appears to be effective in protecting firefighters from toxic chemicals as shown in the contrast with jet-fuel maintenance workers protective gear (as illustrated in figures 3(a) and (b)). Although the overall internal doses experienced from firefighting were shown to be relatively low, future work should focus on the particular events and situations that could give individuals an outlier exposure. Future tests should also be cognizant of the potential for post-work re-exposures, possibly from residual inhalation exposures or dermal exposures once the protective gear is removed.

Acknowledgements

The authors are thankful for the volunteer subjects from the Chicago Fire Department. This research has been subjected to EPA review and approved for publication. This study was internally funded by NIOSH and approved by the NIOSH Human Subjects Review Board. The findings and conclusions in this paper are those of the authors and do not necessarily represent the views of NIOSH.

References

- Amann A 2009 Breath 2009: International conference on breath and breath odor research (26–30 April 2009, Dortmund, Germany) *J. Breath Res.* **4** 010201
- Beauchamp J D and Pleil J D 2013 Simply breath-taking? Developing a strategy for consistent breath sampling *J. Breath Res.* **7** 042001
- Braun P X, Gmachl C F and Dwaek R A 2012 Bridging the collaborative gap: realizing the clinical potential of breath analysis for disease diagnosis and monitoring—tutorial *IEEE Sensors J.* **12** 3258–70
- Cone D C, MacMillan D S, Van Gelder C, Brown D J, Weir S D and Bogucki S 2005 Noninvasive fireground assessment of carboxyhemoglobin levels in firefighters *Prehosp. Emerg. Care* **9** 8–13 (PMID: 16036821)
- Corradi M and Mutti A 2012 News from the Breath Analysis Summit 2011 *J. Breath Res.* **6** 020201
- Davis C E, Epton M, Frank M, Gryshuk A and Kenyon M J 2013 Emerging topics and new developments in the field: the 2012 International Breath Analysis meeting *J. Breath Res.* **7** 039001

- Dunn D H, Devaux I, Stock A and Naeher L P 2009 Application of end-exhaled breath monitoring to assess carbon monoxide exposures of wildland firefighters at prescribed burns *Inhal. Toxicol.* **21** 55–61
- Dweik R A 2011 The great challenge for exhaled breath analysis: embracing complexity, delivering simplicity *J. Breath Res.* **5** 03021
- Fent K W et al 2013 *Evaluation of Dermal Exposure to Polycyclic Aromatic Hydrocarbons in Firefighters Report #2010 0156 3196* (National Institute of Occupational Safety and Health) (www.cdc.gov/niosh/hhe/reports/pdf/2010-0156-3196.pdf)
- Ghio A J, Sobus J R, Pleil J D and Madden M C 2012 Controlled human exposures to diesel exhaust *Swiss Med. Wkly.* **142** w13597
- Hubbard H F, Sobus J R, Pleil J D, Madden M C and Tabacchi S 2009 Application of novel method to measure endogenous VOCs in exhaled breath condensate before and after exposure to diesel exhaust *J. Chromatogr. B* **877** 3652–8
- Kim D, Andersen M E, Chao Y C E, Egeghy P P, Rappaport S M and Nylander-French L A 2007 PBTK modeling demonstrates contribution of dermal and inhalation exposure components to end-exhaled breath concentrations of naphthalene *Environ. Health Perspect.* **115** 894–901
- Lorber M N, Gibb H, Grant L, Pinto J, Pleil J D and Cleverly D 2007 Assessment of inhalation exposures and potential health risks that resulted from the collapse of the World Trade Center towers *Risk Anal.* **27** 1203–21
- Lu S S, Sobus J R, Sallsten G, Albin M, Pleil J D, Gudmundsson A, Madden M C, Strandberg B, Wierzbicka A and Rappaport S M 2014 Exposure to diesel exhaust does not affect levels of PAHs in urine *Biomarkers* **19** 332–9
- Miekisch W, Herbig J and Schubert J K 2012 Data interpretation in breath biomarker research: pitfalls and directions *J. Breath Res.* **6** 036007
- Miranda A L, Martins V, Cascao P, Amorim J H, Valente J, Borrego C, Ferreira A J, Cordeiro C R, Viegas D X and Ottmar R 2012 Wildland smoke exposure values and exhaled breath indicators in firefighters *J. Toxicol. Environ. Health A* **75** 831–43
- NFPA (National Fire Protection Agency) 2007a *NFPA 1971 Standard on Protective Ensembles for Structural Fire Fighting and Proximity Fire Fighting* ed M A Quincy (Quincy, MA: NFPA)
- NFPA (National Fire Protection Agency) 2007b *NFPA 1981 Standard on Open-Circuit Self-Contained Breathing Apparatus (SCBA) for Emergency Services* ed M A Quincy (Quincy, MA: NFPA)
- NFPA (National Fire Protection Agency) 2011 *US Fire Department Profile Through 2010* ed M J Karter and G P Stein (Quincy, MA: NFPA)
- Pleil J D 2008 Role of exhaled breath biomarkers in environmental health science *J. Toxicol. Environ. Health B Crit. Rev.* **11** 613–26
- Pleil J D 2009 Influence of systems biology response and environmental exposure level on between-subject variability in breath and blood biomarkers *Biomarkers* **14** 560–71
- Pleil J D and Hanzel A 2012 Submarines, spacecraft, and exhaled breath *J. Breath Res.* **5** 048005
- Pleil J D and Lindstrom A B 1995 Measurement of volatile organic compounds in exhaled breath as collected in SUMMA canisters *J. Chromatogr. B Biomed. Appl.* **665** 271–9
- Pleil J D and Lindstrom A B 1998 Sample timing and mathematical considerations for modeling breath elimination of volatile organic compounds *Risk Anal.* **18** 585–602
- Pleil J D and Sheldon L S 2011 Adapting concepts from systems biology to develop systems exposure event networks for exposure science research *Biomarkers* **16** 99–105
- Pleil J D and Sobus J R 2013a Estimating lifetime risk from spot biomarker data and intra-class correlation coefficients (ICC) *J. Toxicol. Environ. Health A* **76** 747–66
- Pleil J D and Sobus J R 2013b *Mathematical and Statistical Approaches for Interpreting Exhaled Breath Measurements of Organic Compounds* ed A Amann and D Smith (Amsterdam: Elsevier) chapter 1 (Volatile Biomarkers: Non-Invasive Diagnosis in Physiology and Medicine)
- Pleil J D and Stiegel M A 2013 The evolution of environmental exposure science: using breath-borne biomarkers for ‘discovery’ of the human exposome *Anal. Chem.* **85** 9984–90
- Pleil J D, Funk W E and Rappaport S M 2006 Residual indoor contamination from world trade center rubble fires as indicated by polycyclic aromatic hydrocarbon (PAH) profiles *Environ. Sci. Technol.* **40** 1172–7
- Pleil J D, Miekisch W, Risby T H, Madden M C and Sobus J R 2013 Meeting reports for 2013: recent advances in breath biomarker research *J. Breath Res.* **7** 029001
- Pleil J D, Smith L B and Zelnick S D 2000 Personal exposure to JP-8 jet fuel and exhaust at air force bases *Environ. Health Perspect.* **108** 183–92
- Pleil J D, Stiegel M A, Sobus J R, Liu Q and Madden M C 2011 Observing the human exposome as reflected in breath biomarkers: heat map data interpretation for environmental and intelligence research *J. Breath Res.* **5** 037104
- Pleil J D, Stiegel M A and Sobus J R 2011 Breath biomarkers in environmental health science: exploring patterns in the human exposome *J. Breath Res.* **5** 046005
- Pleil J D, Vette A F, Johnson B A and Rappaport S M 2004 Air levels of polycyclic aromatic hydrocarbons following the World Trade Center disaster *Proc. Natl. Acad. Sci. U. S. A.* **101** 11685–8
- Pleil J D, Vossler T, McClenny W A and Oliver K D 1991 Optimizing sensitivity of selective ion monitoring—GC/MS: analyses for EPA’s TO-14 method *J. Air Waste Manage.* **41** 287–93
- Pleil J D, Williams M and Sobus J R 2012 Chemical safety for sustainability (CSS): biomonitoring in vivo data for distinguishing among adverse, adaptive, and random responses from in vitro toxicology *Toxicol. Lett.* **215** 201–7
- Risby T H and Solga S F 2006 Current status of clinical breath analysis *Appl. Phys. B* **85** 421–6
- Sinues P M, Zenobi R and Kohler M 2013 Analysis of the exhalome: a diagnostic tool for the future *Chest* **144** 746–9
- Sobus J R, Pleil J D, McClean M D, Herrick R F and Rappaport S M 2010 Biomarker variance component estimation for exposure surrogate selection and toxicokinetic inference *Toxicol. Lett.* **199** 247–53
- Sobus J R, Tan Y M, Pleil J D and Sheldon L S 2011 A biomonitoring framework to support exposure and risk assessments *Sci. Total Environ.* **409** 4875–84
- Tan Y M, Sobus J R, Chang D, Goldsmith M, Tornero-Velez R, Pleil J D and Dary C C 2012 Reconstructing human exposures using biomarkers of exposure and other clues *J. Toxicol. Environ. Health B Crit. Rev.* **15** 22–38