Toward Understanding Ultrafine Particle Exposures in Indoor Environments



William W Nazaroff

CEE Department UC Berkeley

Advisory Council Meeting BAAQMD, San Francisco, CA 9 May 2012



http://www.crln.org/files/images/candle_flame_0.jpg



http://yourtreasuredlegacy.com/images/elementary-classroom-2.jpg



http://www.coelhoconstruction.com/air_quality.htm



http://static.howstuffworks.com/gif/gas-vs-electric-cooking-1.jpg

http://blog.aarp.org/shaarpsession/traffic.jpg

Acknowledgments & disclaimer

- UC Berkeley: Seema Bhangar and Nasim Mullen
- Aerosol Dynamics: Susanne Hering & Nathan Kreisberg
- Funding:
 - ARB Contract 05-305
 - Thanks to Peggy Jenkins, Dane Westerdahl, Stephanie Parent
- Thanks to householders and schools for their cooperation.
- Disclaimer:
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Ultrafine particles indoors: Background

- Emerging health concerns about UFP exposure
- New evidence about atmospheric UFP
 - Regional nucleation events
 - Motor vehicles as prominent sources
- Independence of UFP from PM_{2.5}
- Likely, most UFP exposure occurs indoors
- However, little is known about
 - UFP levels indoors
 - Influencing factors

Study goals

- Characterize UFP levels in Northern California
 - Convenience sample of seven houses
 - Convenience sample of six classrooms (4 schools)
- Intensive monitoring in each indoor environment
- Characterize factors that influence levels
- Quantify exposures of occupants
- Apportion exposures to major source categories

Study approach

- Experimental packages (indoor & outdoor)
 - Real-time measurement of UFP and copollutants
 - Temperature & proximity sensors w/ data loggers
 - Occupant questionnaires & site inspections
- Field monitoring campaign
 - 7 houses & 6 classrooms
 - Observational monitoring: ~ 3 days at each site
 - Manipulation experiments at each site
- Extensive interpretive analysis of data

Facilitating technology: WCPC



Array of real-time monitoring instruments

Parameter	Instrument	In1	In2	Out
PN (UFP) level	ME-WCPC (TSI 3781)	✓	✓	 Image: A start of the start of
CO ₂ level	LI-COR 820	√		
CO ₂ level	TSI Q-Trak Plus 8554	 ✓ 		✓
CO level	TSI Q-Trak Plus 8554	 ✓ 		\checkmark
Temperature	TSI Q-Trak Plus 8554	 ✓ 		\checkmark
Relative humidity	TSI Q-Trak Plus 8554	 ✓ 		\checkmark
Ozone level	2B Tech Model 202	 ✓ 		\checkmark
Nitric oxide level	2B Tech Model 400	✓		

• Monitoring: 1-min time resolution; 1.5 m height

QA/QC: Overview

- Ozone, NO, CO, CO₂ monitors calibrated ~ monthly against either reference instrument or standard gases.
- WCPC flow rates routinely checked in field
- Side-by-side monitoring conducted at each site.



Average WCPC side-by-side resultsParameterQMEbQMEcQMEdAverage0.951.021.04Std. dev.0.100.140.14

Slope of readings from instruments QMEb, QMEc, QMEd against reference instrument QMEa

Sample WCPC side-by-side data (Indoor, H0)

Site selection: Houses

- Convenience sample
- All from East Bay area of Northern California
- Source-oriented selection criteria
- Aim for higher than average concentrations, but within normal range



Some characteristics of house sites

ID	City	Y built	V (m ³)	Residents ^a
H0	Oakland	1938	320	2 (M, F)
H1	Oakland	1910	315	4 (M, F, m, m)
H2	Oakland	1949	328	4 (M, F, m, m)
H3	Oakland	1928	200	3 (M, F, m)
H4	Oakland	1904	386	4 (M, F, m, m)
H5	Livermore	1993	420	1 (F)
H6	Emeryville	1996	314	3 (M, M, F)

^a M — male adult, F — female adult, m — male child

House sites: Proximity to major roadways



Some illustrative details: Attributes of H6

- Located in Emeryville, CA
- Built in 1996
- Occupants: 3 adults
- Pilotless gas cooking range
- Used candles one time
- Air-exchange rate (3 measurements): 0.8-0.9 h⁻¹

Site plan at H6

DOWNSTAIRS (Ceiling height= 2.44)



UPSTAIRS (Ceiling height sloped within bedrooms and study)

= sink or

bathtub

= window

PN concentration time series at H6



PN in relation to copollutant data: NO at H6



Cooking activities with gas range or oven: (a), (b), (d), (e), (f), (g), (i), (j); Candle use((c); Toaster oven((h)

Occupancy time-series data at H6



Indoor proportion of outdoor particles at H6

Indoor proportion of outdoor particles

Time (h)	<i>f</i> ₁	<i>f</i> ₂
14.5-18.9	0.54	0.50
25.3-26.4	0.81	0.87
38.5-42.4	0.30	0.28
51.5-54.6	0.33	0.74
62.5-66.4	0.47	0.48
Average	0.44	0.51

Approach: Ratio of average indoor to average outdoor PN level for periods when the house was either unoccupied or all occupants were asleep and there was no evidence of the influence of indoor sources on PN levels.

 f_1 based on ground floor PN f_2 based on upstairs PN level

Characterizing indoor PN sources at H6

ID	Source	Time (h)	σ (10 ¹²)	k+a (h ⁻¹)
а	Stove & rice cooker	4.2-5.4	44	3.6
b	Stove (frying)	8.5-10.8	42	1.5
С	Candle	12.5-13.9	26	1.9
d	Stove (frying)	20.0-23.1	56	1.2
e1	Stove (water)	23.1-23.9	36	2.8
e2	Toaster oven	23.9-26.2	5	1.9
f	Stove (frying)	27.4-29.8	39	1.6
g	Stove & microwave	32.8-34.8	46	1.8
h	Toaster oven	43.4-44.9	35	1.7
i	Stove (water & frying)	47.6-50.3	35	1.5
j	Stove & GF grill	55.6-59.1	40	1.5

 σ = PN emissions (count); k+a = 1st order decay constant

Exposure & apportionment at H6 (3.1 d)

Parameter	R1(F)	R2(M)	R3(M)
Time at home, awake (h)	22.5	28.8	29.8
Time at home, asleep (h)	21.5	28.0	22.5
Time away from home (h)	29.7	16.9	21.4
PN_in1, indoor awake (10 ³ cm ⁻³)	24.7	19.4	23.1
PN_in2, indoor asleep (10 ³ cm ⁻³)	5.3	17.4	7.3
Cumulative exposure (10 ³ cm ⁻³ h)	669	1045	854
Indoor exposure rate (10 ³ cm ⁻³ h/d)	218	341	278
Indoor particles of outdoor origin	23%	26%	29%
Indoor sources, peak events	76%	76%	72%
Indoor, unknown origin	1%	-2%	-1%

Indoor exposure rate = product of average indoor PN concentration $(10^3 \text{ cm}^{-3}) \times \text{occupancy (h/d)}$

All houses: Relationship of PN in to PN out



Indoor PN: Higher when people are awake

Awake at home

Away from home



PN concentration (1000 per cubic centimeter)

Indoor proportion of outdoor particles (f)

• Goal: Determine average indoor concentration of UFP only attributable to average outdoor concentrations.

Site	f ₁	f ₂
H0	0.36	0.37
H1	0.11	
H2	0.51	
H3	0.45	
H4	0.47	0.11
H5	0.29	0.49
H6	0.44	0.51

 Results summary (f₁): avg ± stdev = 0.38 ± 0.14; median = 0.44

Qualitative summary of indoor sources

Source	HO	H1	H2	H3	H4	H5	H6
Gas stove or oven							
Gas clothes dryer							
Furnace (gas fired, central or w all)							
Electric stove (range) or oven							
Toaster or toaster oven							
Ironing clothes							
Candles							
Terpene -based cleaning product use							

- = Reported as not used
- = Used, no clear evidence of emissions
- = Used, individual use associated with an indoor peak
- = Not used or tested alone, joint use with another

potential source associated with an indoor peak

Episodic emissions characterization

- Overall summary: 59 peak events ~ 2.4 events per day
- For peaks associated with distinct activities:
 - Characterized PN emissions (σ) for 40 events
 - Characterized decay constant (k+a) for 38 events

Source	<i>k</i> + <i>a</i> , GM (GSD; N)	σ, GM (GSD; N)
Gas stove	1.8 h ⁻¹ (1.4; 20)	38×10^{12} particles (2.1; 19)
Furnace, central	1.6 h ⁻¹ (1.5; 2)	41 × 10 ¹² particles (1.1; 2)
Candle	1.9 h ⁻¹ (—; 1)	26 × 10 ¹² particles (—; 1)
Toaster oven	1.7 h ⁻¹ (1.2; 4)	9 × 10 ¹² particles (2.8; 4)
Electric stove	1.1 h ⁻¹ (1.3; 5)	10×10^{12} particles (2.1; 4)
Furnace, wall	1.3 h ⁻¹ (1.7; 3)	3.1 × 10 ¹² particles (2.7; 7)
Clothes dryer	2.2 h ⁻¹ (—; 1)	2.2 × 10 ¹² particles (—; 1)
Steam iron	1.5 h ⁻¹ (1.2; 2)	1.9 × 10 ¹² particles (1.4; 2)

PN exposures and apportionment

H0 R1 F adult H0 R2 M adult H1 R1 F adult H1 R2 M adult H1 R3 M child H1 R4 M child H2 R1 F adult H2 R2 M adult H2 R3 M child H2 R4 M child H3 R1 F adult H3 R2 M adult H3 R3 M child H4 R1 F adult H4 R2 M adult H4 R3 M child H4 R4 M child H5 R1 F adult H6 R1 F adult H6 R2 M adult H6 R3 M adult



Occupant

UFP in houses: Key findings

- 1. PN levels in houses were much higher when occupied than when vacant.
- 2. Indoor emission sources are important in study houses.
- 3. Daily average PN exposures per person in houses monitored: $\sim 300 \times 10^3$ cm⁻³ h/d.
- 4. Indoor proportion of outdoor particles in houses monitored: 0.38 ± 0.14 .

Caveats: Small sample of buildings, not statistically representative, few days monitored, one area of California.

Broad extrapolation not warranted!

Site selection: Schools

- Convenience sample
- Elementary schools in the urban portion of the East Bay of Northern California



Sample data: PN concentration vs. time at S1



S1: Occupancy time-series data



S1: Time-average PN levels with occupancy



S1: Source peak from cooking pancakes



S1: PN peak from mopping (manipulation)



Ozone reacts with terpenes in pine oil to form condensable species that first nucleate to form new particles and then condense to cause particle growth.

Summary for classrooms: PN levels



Indoor proportion of outdoor particles (f)

	"Closed"			"Open"		
Site	Time	AER	f ₁ (—)	Time	AER	f ₁ (—)
S1	3%	0.5 h ⁻¹	0.39	96%	2.2 h ⁻¹	0.59
S2	35%	0.4 h ⁻¹	0.16	53%	3.3 h⁻¹	0.54
S3	0%			100%	4.6 h⁻¹	0.76
S4	25%	0.3 h ⁻¹	0.46	68%	3.9 h ⁻¹	0.59
S5	0%			100%	1.9 h ⁻¹	0.51
S6	76%	0.6 h ⁻¹	0.51	17%	4.0 h ⁻¹	0.60
Avg.	23%	0.45 h ⁻¹	0.38	72%	3.1 h ⁻¹	0.60

(*) "Closed" = doors closed and air off; "Open" = door(s) open and/or air on; all data apply for conditions when students were present in classroom.

Summary for classrooms: PN exposure rates





Average ± standard deviation

Students: 50 ± 22 Teachers: 80 ± 40

Units: 10³ cm⁻³ h/d

• Exposure rate is product of average concentration $(cm^{-3}) \times average$ occupancy duration (h/d).

UFP in classrooms: Key findings

- 1. PN levels in classrooms were much higher when occupied than when vacant.
- 2. Indoor emission sources were not important in classrooms.
- Daily average PN exposures per person: students ~ 50 × 10³ cm⁻³ h/d teachers ~ 80 × 10³ cm⁻³ h/d
- 4. Indoor proportion of outdoor particles in classrooms: 0.57 ± 0.10 .
- Caveats: Small sample of buildings, not statistically representative, few days monitored, one area.
- \Rightarrow Broad extrapolation not warranted!

For more information about this study...

NA Mullen, S Bhangar, SV Hering, NM Kreisberg, WW Nazaroff, Ultrafine particle concentrations and exposures in six elementary school classrooms in northern California, *Indoor Air* **21**, 77-87, 2011.



Nasim Mullen



Seema Bhangar

S Bhangar, NA Mullen, SV Hering, NM Kreisberg, WW Nazaroff, Ultrafine particle concentrations and exposures in seven residences in northern California, *Indoor Air* **21**, 132-144, 2011.

Summary remarks on UFP exposure

- High spatial (S) and temporal (T) variation ⇒ Great challenge to use traditional monitoring approaches for characterizing exposure
- Source-oriented perspective
 - Regional nucleation events (T variability dominates)
 - Motor vehicle emissions: time spent in or near traffic (S variability dominates)
 - Indoor sources matter: combustion, high T, ozone + terpenes (S and T variability are both key)
- Importance of source-receptor proximity
- Control opportunities
 - Source reduction
 - Proximity management
 - Air filtration