

1 **Supplementary Material**

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3 **Particle size distribution of nitrated and oxygenated**
4 **polycyclic aromatic hydrocarbons (NPAHs and OPAHs) on**
5 **traffic and suburban sites of a European megacity: Paris**
6 **(France)**

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1 Table S1. Operating characteristics of the MOUDI.

Stage	50 % cut-off diameter (μm)
Inlet	18
1	10
2	5.6
3	3.2
4	1.8
5	1.0
6	0.56
7	0.32
8	0.18
9	0.10
10	0.056
Total filter	0.01*

2 * limit of particle size collected on the total filter has been selected arbitrarily

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4 Changes in flow during and between sample collections would introduce variability in the size
5 of particles collected on each impactor stage. The flow measured at the beginning and end of
6 each run differed by less than 10 %. This variation would cause a change of approximately 5
7 % in the impactor stage cut-off sizes. Because the flow was not recorded during sampling and
8 these sampling errors introduce only minor variations in D_{p50} , for the remainder of this work
9 the air flow is taken to be constant ($1.8 \text{ m}^3 \text{ h}^{-1}$).

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1 Reagents and materials

2 Chemical reagents and gases used in this study are reported in Table S1, together with the
3 name of suppliers and the purity grades.

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5 Table S2. List and characteristics of the chemicals, solvents and gases used.

Compound	Supplier	Purity (%)
<i>OPAHs</i>		
1-Naphthaldehyde	Acros	95
9-Fluorenone	Acros	99
9-Phenanthrenecarboxaldehyde	Aldrich	97
9,10-Anthaquinone	Acros	98
1,4-Anthaquinone	Chiron	97.8
Benzo[a]fluorenone	Chiron	99.9
Benzo[b]fluorenone	Chiron	99.8
Benzanthrone	Acros	99
Benz[a]anthracen-7,12-dione	Acros	99
<i>NPAHs</i>		
1-Nitronaphthalene	Cluzeau	99
2-Nitronaphthalene	Cluzeau	99
2-Nitrofluorene	Cluzeau	98
9-Nitroanthracene	Chiron	86.6
9-Nitrophenanthrene	Cluzeau	100
3-Nitrophenanthrene	Cluzeau	99.7
3-Nitrofluoranthene	Cluzeau	99.5
1-Nitropyrene	Cluzeau	99
2-Nitropyrene	Cluzeau	99.9
4-Nitropyrene	Chiron	99.8
7-Nitrobenz[a]anthracene	Cluzeau	99
6-Nitrochrysene	Cluzeau	98
1,3-Dinitropyrene	Cluzeau	99.9
1,6-Dinitropyrene	Cluzeau	97.5
1,8-Dinitropyrene	Chiron	99.9
1/3-Nitrobenzo[a]pyrene	Chiron	99.7
6-nitrobenzo[a]pyrene	Cluzeau	99.8
<i>Labelled deuterium NPAHs/OPAHs</i>		
3-Nitrofluoranthene-d9	Cluzeau	99.3
1-Nitronaphthalene-d7	Cluzeau	99.4
2-Nitrofluorene-d9	Cluzeau	98.8
6-Nitrochrysene-d11	Cluzeau	99.7
Anthraquinone-d8	Cluzeau	98.6
1-Nitropyrene-d9	Cluzeau	99.2
<i>Solvent</i>		
Methylene Chloride	Sigma-Aldrich	> 99.8
Pentane	Sigma-Aldrich	> 99.0
Isooctane	Sigma-Aldrich	> 99.5
Acetonitrile	VWR	> 99.9
<i>Gases</i>		
Helium	Air Liquide	99.9999
Nitrogen	Air Liquide	99.999
Methane	Air Liquide	99.995
Argon	Air Liquide	99.9999

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1 **Analytical procedures**

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3 **Extractions**

4 Sample extractions were realised using pressurized solvent extraction (Dionex, ASE 200)
5 with CH₂Cl₂ as solvent. Extraction procedure was the following: 33 mL cells at 120 °C, 140
6 bars, 3 cycles of 6 minutes, flush 50 % and purge 120 seconds.

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8 **NPAH and OPAH analyses**

9 Prior to analyses, extracts were purified on solid phase extraction (SPE) (alumina and silica
10 (Upti-clean Aln 500 mg/3 mL, interchim and Upti-clean Si-S 500 mg/3 mL, Interchim)) using
11 a protocol described in a previous publication (Albinet et al., 2006). After purification,
12 extracts were evaporated under argon stream near to dry and dissolved in isooctane.

13 NPAHs and OPAHs were analysed by GC/MS, using a Perkin-Elmer Clarus 500 coupled with
14 a Perkin-Elmer Turbomass gold in the NICI mode in selective ion monitoring mode (Albinet
15 et al., 2006). The column used was a DB-5MS (30 m × 0.25 mm × 0.25 μm film thickness,
16 Agilent J&W). Program settings were as follow: gas flow at 1.2 mL min⁻¹, cool splitless
17 injection (40 to 320 °C) of 1 μL, and transfer line at 300 °C. The initial oven temperature was
18 60 °C for 2 min, then increasing at 45°C min⁻¹ until 150 °C for 5 min; and 5 °C min⁻¹ to 300
19 °C for 7 min. Total run time was 46 min. MS parameters were as follow: electron energy 45
20 eV; source temperature 150 °C; methane was used as the reagent gas for NICI. Monitored
21 ions and deuterated standards were listed in Table S2.

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1 Table S3. Selected ion monitoring conditions for OPAHs and NPAHs.

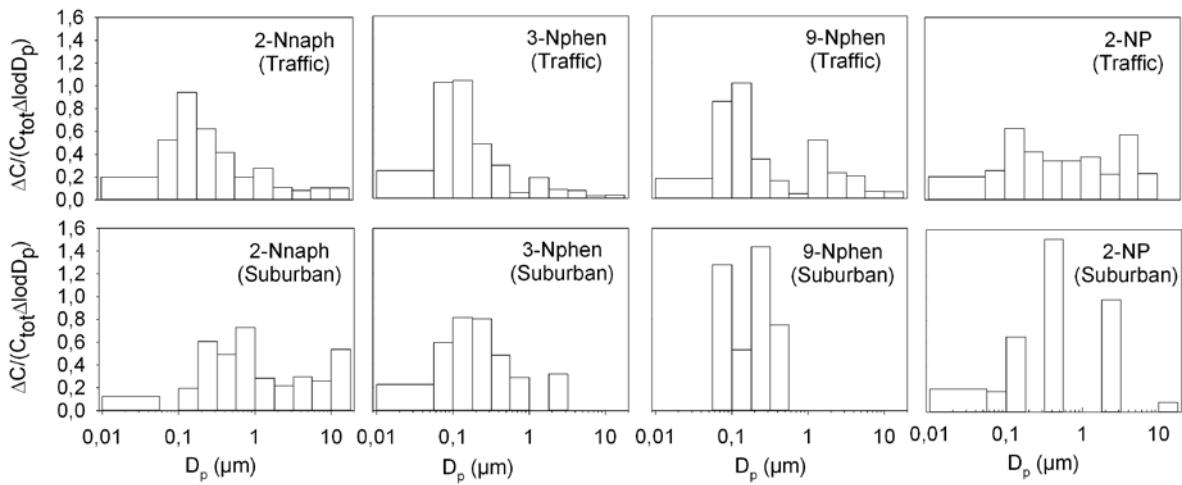
Compounds	Monitored ions (m/z)	Dwell time (s)	Labelled internal standards	Monitored ions (m/z)	Dwell time (s)
<i>OPAHs</i>					
1-Naphthaldehyde	156	0.08	Anthraquinone-d8	216	0.08
9-Fluorenone	180	0.04	Anthraquinone-d8	216	0.08
9-Phenanthrenecarboxaldehyde	206	0.08	Anthraquinone-d8	216	0.08
9,10-Anthraquinone	208	0.04	Anthraquinone-d8	216	0.08
1,4-Anthraquinone	208	0.04	Anthraquinone-d8	216	0.08
Benzo[a]fluorenone	230	0.04	Anthraquinone-d8	216	0.08
Benzo[b]fluorenone	230	0.04	Anthraquinone-d8	216	0.08
Benzanthrone	230	0.04	Anthraquinone-d8	216	0.08
Benz[a]anthracen-7,12-dione	258	0.08	Anthraquinone-d8	216	0.08
<i>NPAHs</i>					
1-Nitronaphthalene	173	0.08	1-Nitronaphthalene-d7	180	0.04
2-Nitronaphthalene	173	0.08	1-Nitronaphthalene-d7	180	0.04
2-Nitrofluorene	211	0.08	2-Nitrofluorene-d9	220	0.08
9-Nitroanthracene	223	0.04	2-Nitrofluorene-d9	220	0.08
9-Nitrophenanthrene	223	0.04	2-Nitrofluorene-d9	220	0.08
3-Nitrophenanthrene	223	0.04	2-Nitrofluorene-d9	220	0.08
2+3-Nitrofluoranthene^a	247	0.08	3-Nitrofluoranthene-d9	256	0.08
1-Nitropyrene	247	0.08	3-Nitrofluoranthene-d9	256	0.08
2-Nitropyrene	247	0.08	3-Nitrofluoranthene-d9	256	0.08
4-Nitropyrene	247	0.08	3-Nitrofluoranthene-d9	256	0.08
1-Nitropyrene-d9	256	0.08	3-Nitrofluoranthene-d9	256	0.08
7-Nitrobenz[a]anthracene	273	0.08	6-Nitrochrysene-d11	284	0.08
6-Nitrochrysene	273	0.08	6-Nitrochrysene-d11	284	0.08
1,3-Dinitropyrene	292	0.04	6-Nitrochrysene-d11	284	0.08
1,6-Dinitropyrene	292	0.04	6-Nitrochrysene-d11	284	0.08
1,8-Dinitropyrene	292	0.04	6-Nitrochrysene-d11	284	0.08
1-Nitrobenzo[a]pyrene	297	0.08	6-Nitrochrysene-d11	284	0.08
3-Nitrobenzo[a]pyrene	297	0.08	6-Nitrochrysene-d11	284	0.08
6-Nitrobenzo[a]pyrene	297	0.08	6-Nitrochrysene-d11	284	0.08

2 ^a The separation of these two isomers could not be achieved on the DB-5MS column.

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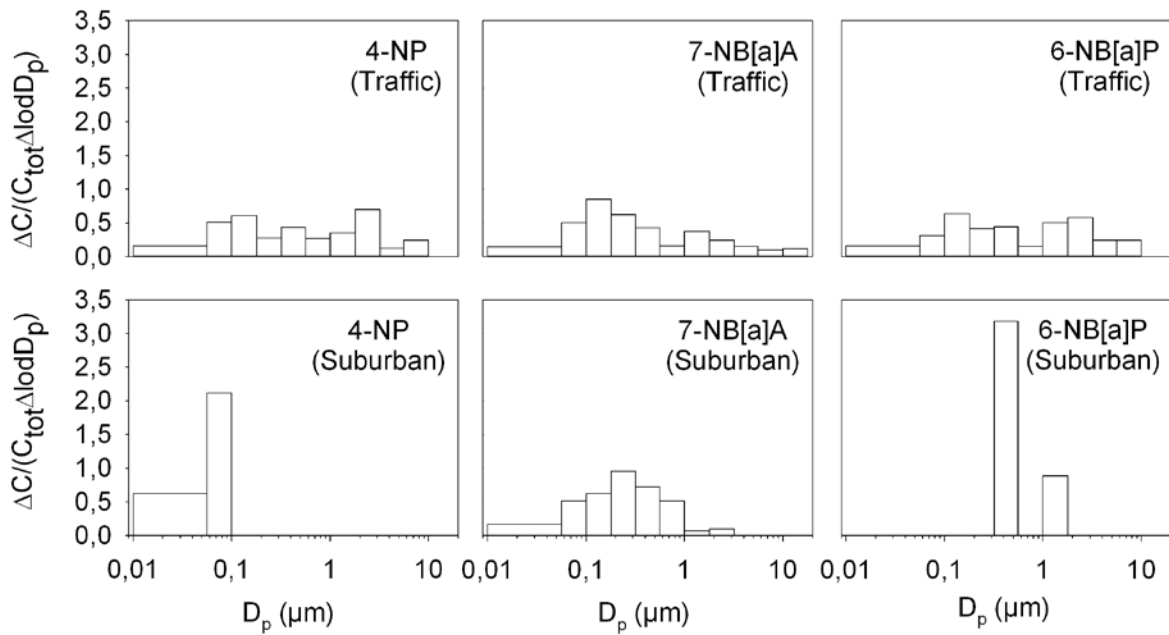
1 **Additional figures**



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3 Figure. S1. Particle size distributions of 2-nitronaphthalene (2-Nnaph), 9-nitrophenanthrene
4 (9-Nphen), 3-nitrophenanthrene (3-Nphen) and 2-nitropyrene (2-NP) at the traffic and
5 suburban sites.

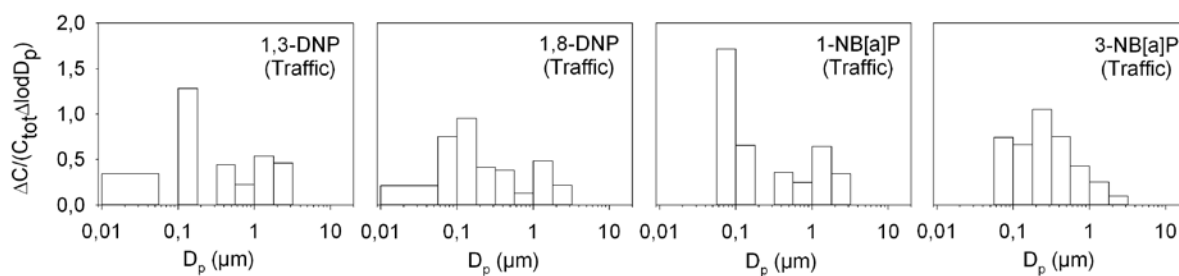
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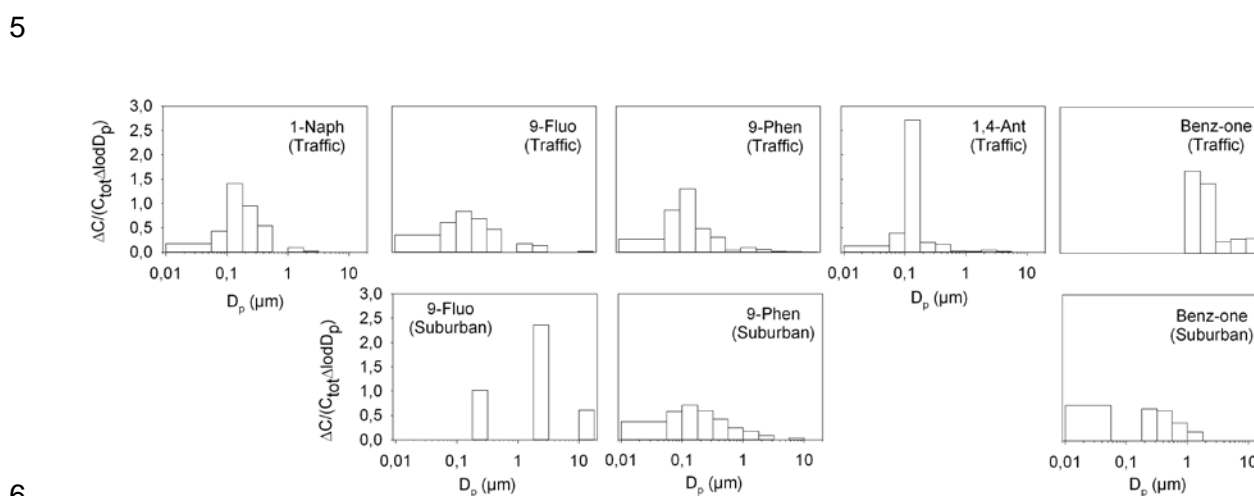
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8 Figure. S2. Particle size distributions of 4-nitropyrene (4-NP), 7-nitrobenz[a]anthracene (7-
9 NB[a]A) and 6-nitrobenzo[a]pyrene (6-NB[a]P) at the traffic and suburban sites.

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2 Figure. S3. Particle size distributions of 1,3-dintropyrene (1,3-DNP), 1,8-dintropyrene (1,8-
3 DNP), 1-nitrobenzo[a]pyrene (1-NB[a]P) and 3-nitrobenzo[a]pyrene (3-NB[a]P) at the traffic
4 suburban sites. Missing graphs correspond to the compounds not quantified.



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7 Figure. S4. Particle size distributions of 1-naphthaldehyde (1-Naph), 9-fluorenone (9-Fluo), 9-
8 phenanthrenecarboxaldehyde (9-Phen), 1,4-anthraquinone (1,4-Ant) and benzanthrone (Benz-
9 one) at the traffic and suburban sites. Missing graphs correspond to the compounds not
10 quantified.

11
12 **References**

13 Albinet, A., Leoz-Garziandia, E., Budzinski, H., and Villenave, E.: Simultaneous analysis of
14 oxygenated and nitrated polycyclic aromatic hydrocarbons on standard reference material
15 1649a (urban dust) and on natural ambient air samples by gas chromatography-mass
16 spectrometry with negative ion chemical ionisation, Journal of Chromatography A, 1121,
17 106-113, 2006.