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1        **Euro 6 unregulated pollutant characterization and statistical analysis of**  
2        **the impact of aftertreatment devices and driving conditions on recent**  
3        **passenger cars emissions**

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10

11        **Abstract**

12        This study aims to measure and analyze unregulated compound emissions for two Euro 6 diesel and  
13        gasoline vehicles. The vehicles were tested on a chassis dynamometer under various driving cycles:  
14        Artemis driving cycles (urban, road and motorway), the New European Driving Cycle (NEDC) and the  
15        World Harmonized Light-Duty Test Cycle (WLTC) for Europe, and world approval cycles. The  
16        emissions of unregulated compounds — such as total particle number (PN) (over 5.6 nm), black  
17        carbon (BC), NO<sub>2</sub>, BTEX (benzene, toluene, ethylbenzene and xylene), carbonyl compounds and  
18        polycyclic aromatic hydrocarbons (PAHs) — were measured with several on-line devices and  
19        different samples were collected using cartridges and quartz filters. Furthermore, a preliminary  
20        statistical analysis was performed on eight Euro 4-6 diesel and gasoline vehicles to study the impacts  
21        of driving conditions and aftertreatment and engine technologies on emissions of regulated and  
22        unregulated pollutants. The results indicate that urban conditions with cold start induce high  
23        emissions of BTEX and carbonyl compounds. Motorway conditions are characterized by high  
24        emissions of particle numbers and CO, which mainly induced by gasoline vehicles. Compared with  
25        gasoline vehicles, diesel vehicles equipped with catalyzed or additive DPF emit fewer particles but  
26        more NO<sub>x</sub> and carbonyl compounds.

27

28        **Keywords**

29        Regulated and unregulated pollutants; Emission factors; Euro 6 vehicles; Chassis dynamometer;  
30        Driving conditions; Aftertreatment systems

31

32

33 **Abbreviation list**

34

35	DPF	Diesel Particulate Filter
36	DI	Direct Injection
37	NEDC	New European Driving Cycle
38	WLTC	World harmonize Light-duty Test Cycle
39	PN	Particle Number
40	BC	Black Carbon
41	BTEX	Benzene, Toluene, Ethylbenzene, Xylene
42	PAH	Polycyclic Aromatic Hydrocarbon
43	DOC	Diesel Oxidation Catalyst
44	SCR	Selective Catalytic Reduction
45	TWC	Three-Way Catalyst
46	PCA	Principal Component Analysis
47	DCI	Direct Common rail Injection
48	CVS	Constant Volume Sampler
49	CPC	Condensation Particle Counter
50	PMP	Particle Measurement Program
51	ELPI	Electrical Low Pressure Impact
52	FMPS	Fast Mobility Particle Sizer
53	AE	Aethalometer
54	ArtUrb C	Artemis Urban with Cold start
55	ArtUrb H	Artemis Urban with Hot start
56	ArtROAD	Artemis Rural
57	ArtMW	Artemis Motorway
58	Add DPF	Additive DPF
59	Cat DPF	Catalyzed DPF

60

## 61 Introduction

62 Road transportation (more particularly, light-duty vehicles) is one of the main causes of air  
63 pollution. In urban areas, road traffic represents the main source of emissions of regulated pollutants  
64 as well as unregulated pollutants, such as BTEX, PAHs, and carbonyl compounds (1). Several of these  
65 pollutants have an important role in climate change while others could lead to serious negative  
66 impacts on human health (2-4).

67 To reduce road traffic emissions, the European Union is imposing increasingly stringent emission  
68 limits for regulated compounds. Various aftertreatment devices — such as the diesel oxidation  
69 catalyst (DOC), the diesel particulate filter (DPF), the selective catalytic reduction (SCR) or NO<sub>x</sub> trap  
70 and the three-way catalyst (TWC) — are being used to bring the pollutant emissions below  
71 regulatory levels (5-7). Although these technologies make it possible to significantly reduce regulated  
72 compound emissions, they affect some emissions of pollutants. Catalyzed or additive DPF reduce  
73 particle mass emission, with efficiency near 100%, but they might induce an increase of fine and  
74 ultrafine particle emissions, and affect NO<sub>2</sub>, volatile organic compound, PAH, BTEX, and black carbon  
75 (BC) emissions (8-15). The actual impacts of these aftertreatment technologies on unregulated  
76 pollutant emissions are not fully known (13, 16-20). The recent study by Louis et al. (14) showed that  
77 catalysed DPF vehicles emitted about 3 to 10 times more carbonyl compounds and particles than  
78 additive DPF vehicles, respectively, during urban driving cycles, while additive DPF vehicles emitted 2  
79 and 5 times more BTEX and carbonyl compounds during motorway driving cycles.

80 Vehicle emissions are also affected by driving conditions. In the case of diesel vehicles, urban  
81 driving conditions or high engine RPM (revolutions per minute) involves high emissions of CO, NO<sub>x</sub>  
82 and HC compared to a steady speed profile or low engine speed (21-23). For gasoline vehicles, cold  
83 start and high-speed conditions induce high emissions of the same compounds (21-25). For urban  
84 driving conditions, diesel vehicles emit more NO<sub>x</sub> during hot start compared to cold start (13, 14).  
85 Various studies also show that cold start results in significant emissions of BC, PAHs, BTEX and  
86 aldehyde (9, 12, 14, 21, 24-28) compared to urban hot start. A vehicle running low speed produces  
87 high PAH and carbonyl compound emissions while a vehicle running at high speed produces high  
88 particulate emissions (12, 26-28). Emission factors of unregulated compounds reported in the  
89 aforementioned papers have been measured with Euro 1 to Euro 5 diesel and gasoline vehicles with  
90 a relatively small number of vehicle samples: six Euro 4–5 vehicles by Louis et al. (15), four Euro 2–4  
91 vehicles by Rehn (25), and 25 Euro 1–3 vehicles by Caplain et al. (26). Emissions of unregulated  
92 compounds by Euro 6 vehicles have not been yet measured. Moreover, unregulated compound  
93 emission factors were often measured under Artemis driving conditions. Impacts of the WLTC (World  
94 Harmonize Light-Duty Test Cycle, future world approved driving cycle) on such emissions have not  
95 been studied to date.

96 In this paper, two recent in-use Euro 6 vehicles – diesel with catalyzed DPF and NO<sub>x</sub> trap and  
97 gasoline direct injection (DI) with propulsion engine – were tested under Artemis urban, road and  
98 motorway, WLTC, and NEDC (New European Driving Cycle) driving cycles. Unregulated compound  
99 emissions were measured to improve knowledge on their emissions under different driving  
100 conditions and supplement the emission factor database used by diverse emission models.  
101 Furthermore, Clairotte et al. used the Principal Component Analysis (PCA) statistical analysis method  
102 for two mopeds (31) and two light duty flexible-fuel vehicles (19) to investigate the impact of driving  
103 conditions on vehicle emissions. This method makes it possible to analyze the effects of low  
104 temperature on cold start gaseous emissions and deeply characterize online emission patterns. Our

105 research thus used preliminary PCA to study the impacts of technologies and driving conditions on  
 106 regulated and unregulated emissions of pollutants from eight Euro 4–6 diesel and gasoline vehicles  
 107 tested in this research and previous studies (15).

## 108 1. Materials and Methods

### 109 1.1.Characteristics of the vehicles

110 One Euro 6 gasoline vehicle with a direct injection system, TWC, and propulsion engine (Vehicle  
 111 1) and one Euro 6 diesel vehicle with DOC, catalyzed DPF, and NO<sub>x</sub> trap (Vehicle 2) were tested. The  
 112 propulsion engine was located at the rear of the rear-wheel drive vehicle. The technical  
 113 characteristics of the two Euro 6 vehicles are given in Table 1.

114 **Table 1. Technical characteristics of tested Euro 6 diesel and gasoline vehicles**

Vehicle	No. 1	No 2
Size class	0.9 DI	1.5 DCI
Technology	Gasoline	Diesel
Standard	Euro 6b	Euro 6b
Engine capacity (cm <sup>3</sup> )	999	1461
Empty weight (kg)	864	1087
Mileage (km)	2164	4700
Gearbox type	Manual (5)	Manual (5)
Aftertreatment systems	TWC	DOC + Catalyzed DPF + NO <sub>x</sub> trap
Registration date	12/11/2015	12/31/2015
Test date	03/31/2016	04/14/2016

### 115 1.2.Experimental set-up

116 Vehicle emissions were measured on the chassis dynamometer at the Transport and  
 117 Environment Laboratory (LTE) of the French Institute of Science and Technology for Transport,  
 118 Development and Networks (IFSTTAR). Exhaust gas was collected at the outlet of the tailpipe, and  
 119 sent through the constant volume sampler (CVS) to be diluted with filtered air. Pollutant emissions  
 120 were measured at the outlet of the CVS by various on-line gas and particulate analyzers and were  
 121 sampled on various filters and cartridges for off-line analysis.

122 Measurements were performed for five different driving cycles — NEDC, European approval  
 123 cycle, WLTC, world approval cycle, and Artemis urban, road and motorway driving — cycles that are  
 124 more representative of real-world driving conditions (29, 30). The characteristics of the driving cycles  
 125 are given in Table 2. Each cycle was repeated two to six times under the same experimental  
 126 conditions in order to check the emission level and exclude high emission vehicles. The only  
 127 exception was NEDC, for which only one measurement was conducted for each vehicle. All the  
 128 experiments (this research and previous studies) were performed using commercial fuel (less than 10  
 129 ppm sulfur content) from the same filling station to minimize the impact of fuel composition on  
 130 emissions. All the diesel and gasoline vehicles were filled with fuel meeting the requirements of EN

131 590 and EN 228, respectively. The detailed fuel properties are given in the Table S1 in the Supporting  
132 Information.

133 **Table 2. Characteristics of the driving cycles**

Cycles	Start	Mileage (km)	Duration (s)	Average speed (km/h)	CVS flow (m <sup>3</sup> /min)
NEDC	Cold	11.2	1180	34	9
WLTC	Cold	23.25	1800	47	13
Artemis Urban	Cold/Hot	4.47	921	17	9
Artemis Road	Hot	14.7	862	61	9
Artemis Motorway	Hot	23.7	736	116	13

### 134 1.3. Analytical methods

135 Concentrations of regulated compounds used in PCA analysis (Section 3.4 herein) were measured  
136 using a HORIBA analytical emissions system. The analyzer using infrared absorption principle was  
137 used to measure carbon monoxide (CO), carbon dioxide (CO<sub>2</sub>), flame ionization detection to total  
138 hydrocarbon (THC) and methane (CH<sub>4</sub>) and chemiluminescence for nitrogen oxides (NO<sub>x</sub>) and  
139 nitrogen oxide (NO). The concentration of nitrogen dioxide (NO<sub>2</sub>) is determined by subtracting NO  
140 from NO<sub>x</sub>. The concentration of CO<sub>2</sub> was also measured with a MIR-2M (Multi-gas InfraRed;  
141 Environment SA), which also uses infrared absorption. Both CO<sub>2</sub> analyzers show good correlation  
142 between measurements, with a relative gap of about 2%.

143 The total particle number was measured with condensation particle counter (CPC, 3775 TSI). The  
144 CPC has a butanol condensation chamber enabling the detection of particles between 4 nm and 2  
145 μm. The instrument was operated once a second at 1.5 L/min, with a concentration range of 0 to 10<sup>7</sup>  
146 particle/cm<sup>3</sup>. The Particle Measurement Program (PMP) has been proposed by the direction of Joint  
147 Research Center (JRC), a Directorate-General of the European Commission, as regulatory method for  
148 measuring particle numbers for Euro 5 and Euro 6 vehicles. PMP makes it possible to remove volatile  
149 particles with a 50% cut-point size of 23 nm. One of the main reasons for cutting volatile particles is  
150 that the measurement of non-volatile particles is more repeatable. However, studies by Louis et al.  
151 (15) showed that most particles emitted by tested Euro 4–5 vehicles were ultrafine particles with  
152 diameters of less than 23 nm. In order to obtain the fullest amount of data on total particle number  
153 emissions, the PMP was not used in this study. Furthermore, taking into account this volatile part,  
154 standard variations of particle number quantification with six repeated driving cycles ranged  
155 between 7 and 20%, which was quite low.

156 The particle size number distribution was measured with two different devices. The first was the  
157 Electrical Low Pressure Impactor (ELPI; DEKATI), which has 12 filter stages and determines particle  
158 number distributions from 7 nm to 10 μm. The ELPI was operated once a second at a flow rate of 10  
159 L/min. The minimal detection limit ranged from 250 to 0.1 particles/cm<sup>3</sup> depending on the impactor  
160 stage. The second device was the Fast Mobility Particle Sizer (FMPS; TSI), which measures the total  
161 particle number and distribution ranging from 5.6 to 560 nm, with a concentration range from 0 to  
162 10<sup>7</sup> particle/cm<sup>3</sup> and with a flow rate of 8 L/min. The particle numbers obtained by these three  
163 devices were fairly well correlated, with a relative gap of about 20%. This gap might be explained by  
164 the fact that measurements of the size range of particles are not the same for all three devices.

165 The black carbon concentration was measured using an aethalometer (AE 33-7, Magee  
166 Scientific). The experimental data were collected once a second with the instrument operating at a  
167 flow rate of 5 L/min. The detection limit for 1 hour was 5 ng/m<sup>3</sup> with a concentration range of 10 to  
168 10<sup>5</sup> ng/m<sup>3</sup>. Light attenuation was measured at seven wavelengths, from UV to IR (370, 470, 525, 590,  
169 660, 880 and 940 nm). The 880 nm wavelength corresponding to the maximum amount of black  
170 carbon was used for black carbon quantification in this study.

171 Unregulated compounds, such as BTEX, carbonyl compounds and PAHs, were sampled on various  
172 cartridges or quartz filters. Emissions of three repeated Artemis cycles or two repeated WLTC cycles  
173 were sampled on one cartridge to collect enough pollutants for chemical analysis. Before each  
174 exhaust sample, one blank sample (dilution air in CVS) was collected under the same experimental  
175 conditions. Gas phase PAHs were collected with ORBO 43 cartridges. The sampling flow rate was 0.5  
176 L/min. Particulate phase PAHs were collected on quartz filters at a flow rate of 50 L/min. BTEX were  
177 collected on Tenax cartridges at a flow rate of 0.5 L/min. Carbonyl compounds were collected on  
178 DNPH cartridges at a flow rate of 2 L/min. The cartridges and filters were analyzed by TERA-  
179 Environment, a private laboratory with standardized analytical methods (for detailed of the analytical  
180 methods see ISO-16000-6, ISO 16000-3, NIOSH 2549, NIOSH 5506 and NF X43-025) (See Table S2 in  
181 the Supporting Information). The complete list of compounds analysed using the cartridges and  
182 filters, which includes six BTEX, 11 carbonyl compounds and 16 PAHs, is given below.

183

- 184 • BTEX: benzene, toluene, ethylbenzene, m-xylene, p-xylene, o-xylene
- 185 • Carbonyl compounds: formaldehyde, acetaldehyde, acetone, acrolein, propionaldehyde,  
186 crotonaldehyde, methacrolein, butanal, benzaldehyde, pantanal, hexanal
- 187 • PAHs (gas and particulate phases): naphthalene, acenaphthylene, acenaphthene, fluorene,  
188 phenanthrene, anthracene, fluoranthene, pyrene, benzo(a)anthracene, chrysene,  
189 benzo(b,j)fluoranthene, benzo(k)fluoranthene, benzo(a)pyrene, dibenz(a,h)anthracene,  
190 benzo(g,h,i)perylene, indeno(1,2,3-c,d)pyrene

#### 191 **1.4. Statistical analysis**

192 The impacts of driving conditions and technologies on emissions produced by two Euro 6 vehicles  
193 (tested in this study) and six Euro 4–5 vehicles (tested in previous studies by Louis et al. (15)) were  
194 investigated using a statistical analysis method, i.e., the Principal Component Analysis (PCA). The  
195 basic objective of PCA is to reduce the data set and find the best space to project the variables. This  
196 dimension reduction process creates a limited number of eigenvectors called “principal  
197 components”, which are linear combinations and explain most of the total variance of the data set.  
198 This method was used by Clairotte et al. (31) to separately analyze two motorcycles with around 20  
199 measured pollutants. In this case, the variables corresponded to emission factors measured for 32  
200 pollutants, cited above, and the individuals were the tested vehicles and driving cycles (Artemis  
201 urban with cold and hot start, road and motorway, NEDC). The SPAD8 software was used to create  
202 geometric interpretations between the variables and individuals from principal components. These  
203 representations allow make it possible to visually restore the relationships between the vehicles, or  
204 the driving cycles, and the emission factors (33).

205 Geometric representations are produced in 2D graphic form. Each dimension corresponds to one  
206 principal component representing the maximum percentage of the total variance for the variable set.  
207 Both dimensions are represented by two factorial axes. The projection of the variables according to  
208 the two dimensions produces a graphic representation called “correlation circle”. In this circle, each  
209 arrow represents a variable (an emission factor in the present case). An arrow close to the circle  
210 means that the circle can be interpreted. Any arrows that are clustered together are correlated and  
211 represent similar emission behavior. For projection of the individuals, the individuals in our case are  
212 vehicles or driving cycles, the PCA provides a factorial plan according to the same two factorial axes.  
213 In this plan, any individuals that are clustered together have closed variable values indicating that  
214 they have similar effects on emissions. This method makes it possible to study the impacts of vehicle  
215 technologies and driving conditions on emission factors.

## 216 **2. Results and Discussion**

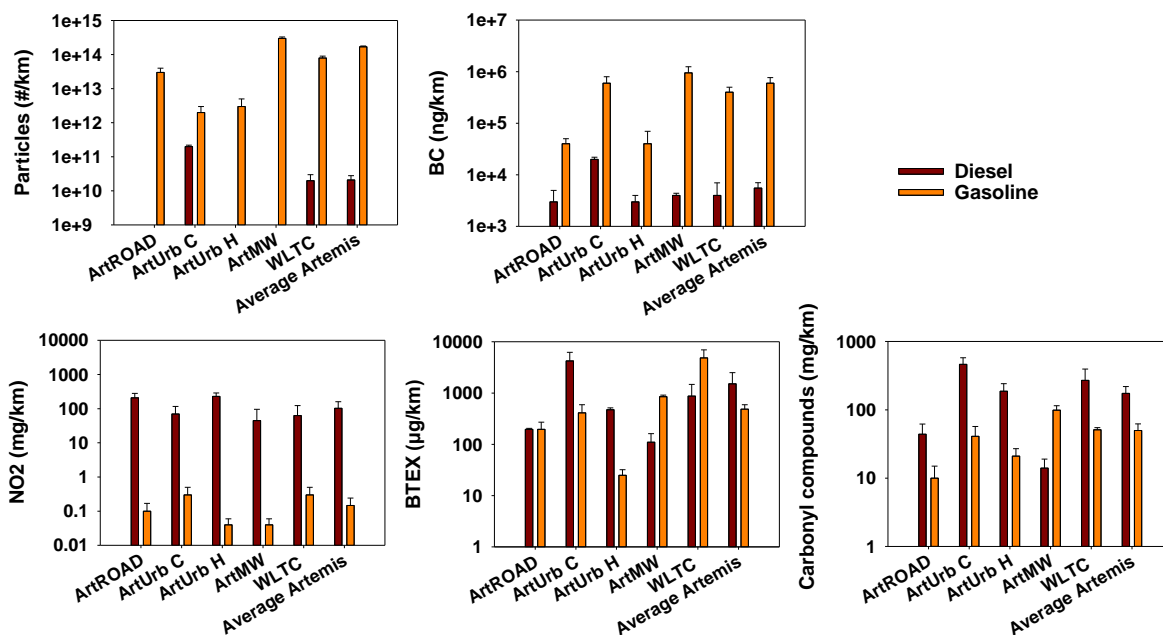
### 217 **2.1. Euro 6 gasoline and diesel emission factors**

218 Particles, BC, NO<sub>2</sub>, BTEX and carbonyl compound emission factors for Euro 6, gasoline and diesel,  
219 vehicles with the six driving conditions (Artemis urban cold start (ArtUrb C), Artemis urban hot start  
220 (ArtUrb H), Artemis road (ArtROAD), Artemis motorway (ArtMW), WLTC) are presented in the Figure  
221 1.

222 The article number emission factors for the gasoline vehicle varied between  $2.3 \times 10^{12}$  to  $2.9 \times 10^{14}$   
223 #/km. The highest factors were obtained for the motorway cycle and the lowest were obtained for  
224 the urban cycle with similar emissions between hot start and cold start. Compared to the diesel  
225 vehicle equipped with DPF to emit few particles, the gasoline vehicle emitted 4 to 4500 times more  
226 particles. The gasoline vehicle emitted 10 to 30 times more BC than the diesel vehicle under Artemis  
227 urban and road driving conditions. Under the Artemis motorway and WLTC with high-speed  
228 conditions, the gasoline vehicle emitted 200 to 250 times more BC than the diesel vehicle. NO<sub>2</sub>  
229 emissions from the gasoline vehicle were low, i.e., between 0.04 and 0.3 mg/km, and were 200 to  
230 5000 less than for the diesel vehicle (45 to 229 mg/km) depending on the driving conditions. For the  
231 unregulated compounds, the BTEX emission factors ranged from 0.03 and 4.9 mg/km for the gasoline  
232 vehicle and 0.11 and 4.2 mg/km for the diesel vehicle. For the exhaust samples, only formaldehyde,  
233 acetaldehyde and acetone were above the quantification limit that could be quantified. The diesel  
234 vehicle emitted on average 11 times more carbonyl compounds than the gasoline vehicle for all the  
235 cycles. The only exception was the motorway cycle, for which the gasoline vehicle emitted seven  
236 times more carbonyl compounds than the diesel vehicle. The concentrations of the sixteen PAHs in  
237 the particulate phase were below the detection limit for both vehicles (See Table S3 in the  
238 Supporting Information).

239 Generally, Artemis urban cold start induces more emissions than hot start for all the pollutants  
240 measured for both the diesel and gasoline vehicles. The diesel vehicle emitted 100 times more PN  
241 under urban cold start condition than hot start. Cold start also induced 9–16, 2.5 and 10 times more  
242 BTEX, carbonyl compound and BC emissions, respectively, than with hot start for the diesel and  
243 gasoline vehicles. However, the diesel vehicle emitted three times more NO<sub>2</sub> under urban hot start  
244 than under cold start. This same emission characteristic has been observed in other studies as well  
245 (13, 14). The high NO<sub>2</sub> emissions in urban driving conditions and the increase in the number of Euro 6  
246 diesel vehicles in the fleet raise serious concerns about urban air quality.





247  
 248 **Figure 1.** Particles, BC, NO<sub>2</sub>, BTEX and carbonyl compounds emission factors for Euro 6 diesel and  
 249 gasoline vehicles.

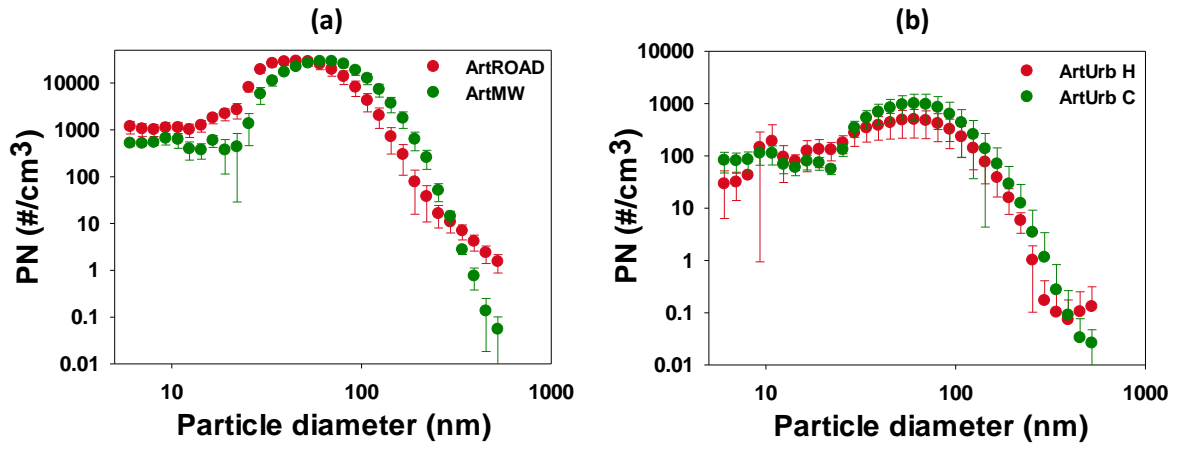
250 Compared to the WLTC approval cycle, diesel emission levels of PN, BC, NO<sub>2</sub>, BTEX and carbonyl  
 251 compounds are similar under the Artemis average cycles (urban cold start + road + motorway). For  
 252 the gasoline vehicle, all the pollutant emissions are similar between the average Artemis and WLTC  
 253 cycles. The only exception is BTEX: the WLTC cycle induced 4 to 13 times more emissions than  
 254 average Artemis cycle. However, only two vehicles in this study were tested using the WLTC cycle.  
 255 Further testing will have to be conducted to confirm the impact of the WLTC on emissions of  
 256 unregulated pollutants.

## 257 2.2. Size distribution of the particle number

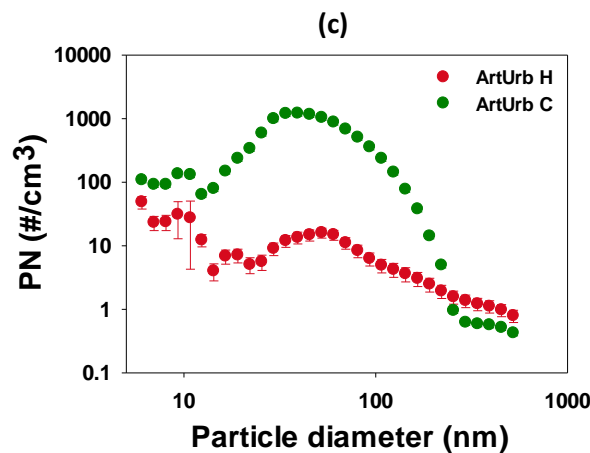
258 The particle size distributions were measured with the FMPS for all the Artemis driving cycles.  
 259 Figure 2 (a) and (b) shows the particle size distributions for the gasoline vehicle under motorway and  
 260 road conditions (Figure 2a) and urban conditions with hot and cold start (Figure 2b). The particle size  
 261 ranged from  $22 \pm 1.4$  to  $220 \pm 15.6$  nm for all the Artemis driving cycles. The peak number  
 262 concentrations were around  $45 \pm 3.6$  nm,  $70 \pm 4.7$  nm and  $60 \pm 4.7$  nm for the road, motorway and  
 263 urban driving conditions, respectively.

264 The particle emissions for the diesel vehicle were near the background level (i.e., undetectable)  
 265 for Artemis road, motorway and urban with hot start (See Table S3 in the Supporting Information).  
 266 Only Artemis urban cold and hot start is presented in Figure 2c. For urban cycle with cold start, the  
 267 particle sizes varied between  $22 \pm 1.4$  to  $220 \pm 15.6$  nm, with two modes around  $10 \pm 0.8$  and  $40 \pm 2.5$   
 268 nm. The results also show that the particles emitted by the diesel vehicle were smaller than those  
 269 emitted by the gasoline vehicle.

270



271  
272

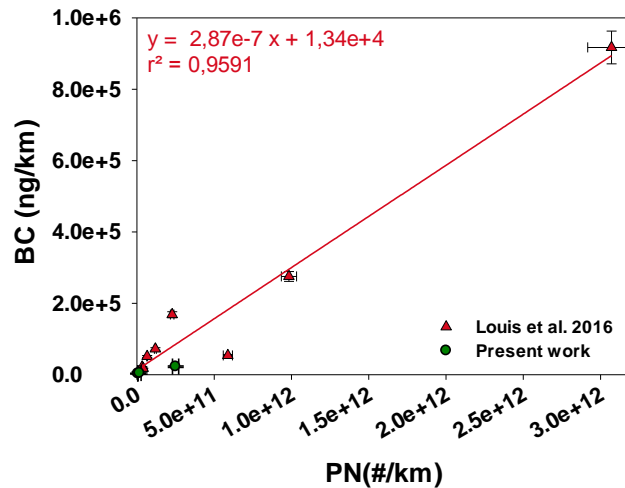


273

274 **Figure 2.** Particle size number distribution for the Euro 6 gasoline DI (a) and (b) and Euro 6 diesel  
275 vehicles (c). (a) The Artemis road (ArtROAD) and Artemis motorway (ArtMW) driving cycles; (b) and  
276 (c) the Artemis urban, hot and cold start (ArtUrb H/C) driving cycles.

### 277 2.3. Pollutant correlations

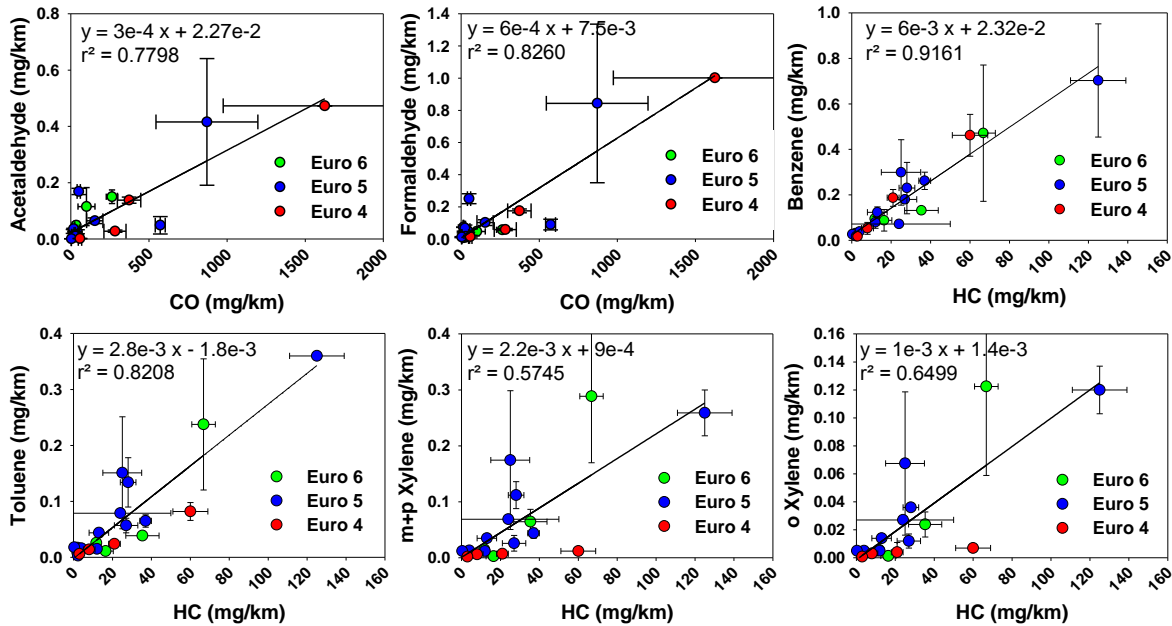
278 The correlations between the pollutant emissions for the two Euro 6 vehicles tested in this study  
279 and the six Euro 4–5 vehicles tested in the previous research (15) were studied. However, only the  
280 emission factors of the diesel vehicles were used because the three gasoline vehicles did not show  
281 good correlations between their pollutant emissions. This is particularly true for the Euro 6 gasoline  
282 DI vehicle tested in this study and which showed very different emission behaviors. These differences  
283 in behavior may be explained by the different vehicle technologies. The Euro 6 gasoline vehicle has a  
284 propulsion engine that leads to a high exhaust temperature at the outlet of the tailpipe (up to 600 °C  
285 during the motorway phases). The Euro 4 and Euro 5 gasoline vehicles tested by Louis et al. (15) were  
286 fitted with a traction engine that led to a lower exhaust temperature (around 200 °C at the tailpipe).  
287 Ghazikhani et al. (34) showed that the increase in exhaust temperature leads to an increase in  
288 emissions of pollutant such as CO and HC.



289

290 **Figure 3.** BC/PN correlation for the Euro 6 diesel vehicle (green dots, present study) and the Euro 4  
 291 and 5 diesel vehicles (red dots) (15) for emissions measured under the Artemis urban cold start,  
 292 WLTC and NEDC driving cycles.

293 At first, the correlations between PN and BC emissions for the diesel vehicles were studied.  
 294 Figure 3 shows the PN/BC correlation obtained from the Euro 6 diesel vehicle measured in this study  
 295 (green dots) and the correlation obtained by Louis et al. (15) for the Euro 4 (red dots) and 5 diesel  
 296 vehicles (green dots). The PN/BC correlation of the Euro 6 diesel vehicle tested in this study follows a  
 297 similar tendency, compared to the Euro 4 and 5 vehicles for the Artemis urban cold start, WLTC, and  
 298 NEDC driving cycles. However, the PN and BC emissions for the Euro 6 diesel vehicle were low,  
 299 making it difficult to arrive at a clear conclusion with uncertainties.



300

301

302 **Figure 4.** Correlations between the various pollutants measured for the Euro 4 (red dots), Euro 5  
 303 (blue dots) and Euro 6 (green dots) diesel vehicles with all Artemis, WLTC and NEDC driving cycles.

304 The correlations between HC and CO (regulated compounds) and unregulated pollutant  
305 emissions measured for the Euro 6 diesel vehicle (present study) and Euro 4–5 diesel vehicles (15)  
306 were also studied under the Artemis urban, cold and hot start, Artemis road, Artemis motorway,  
307 WLTC, and NEDC driving cycles. Figure 4 shows the correlations between unregulated pollutants  
308 (benzene, toluene, xylene, formaldehyde and acetaldehyde), and regulated pollutants (HC and CO).  
309 The correlations are a positive linear correlation with  $r^2$  varying from 0.57 to 0.91.

310 Unlike regulated compounds, emissions of unregulated compounds are not always measured on  
311 a chassis dynamometer. Based on the correlation equations given in Figure 4, it is therefore possible  
312 to estimate the emission factors of benzene, toluene, xylene, formaldehyde and acetaldehyde with  
313 HC and CO measurements for the Euro 4–6 diesel vehicles. However, these correlations only give an  
314 approximate estimation that takes into account the large standard deviation due to the high  
315 variability of emissions of regulated and unregulated compounds during cold start.

## 316 **2.4. Preliminary PCA analysis**

317 Principal component analysis (PCA) is performed as a preliminary method to study the impacts of  
318 driving conditions, aftertreatment, and engine technologies on pollutant emissions by analyzing the  
319 variable main trends. PCA was performed with emission factors of various gaseous and particulate  
320 pollutants measured in this study for two Euro 6 diesel and gasoline vehicles, as well as for four Euro  
321 5 vehicles and two Euro 4 vehicles measured in previous study (15). The technical characteristics of  
322 these eight vehicles are given in the Table S4 in the Supporting Information. To build the PCA, we  
323 used either the five driving conditions (Artemis urban with cold start, urban with hot start, road,  
324 motorway and NEDC) or the eight tested vehicles as individuals, with the 32 pollutant emissions for  
325 each case (as variables) (Section 2.4 herein). The SPAD8 software was then used to perform the PCA  
326 analysis of the data set. The results showed that the first two dimensions set by PCA account for the  
327 bulk of the total variance. Therefore, in these two cases, only two dimensions were used for this  
328 preliminary study.

### 329 **2.4.1. Impacts of driving conditions on pollutant emissions**

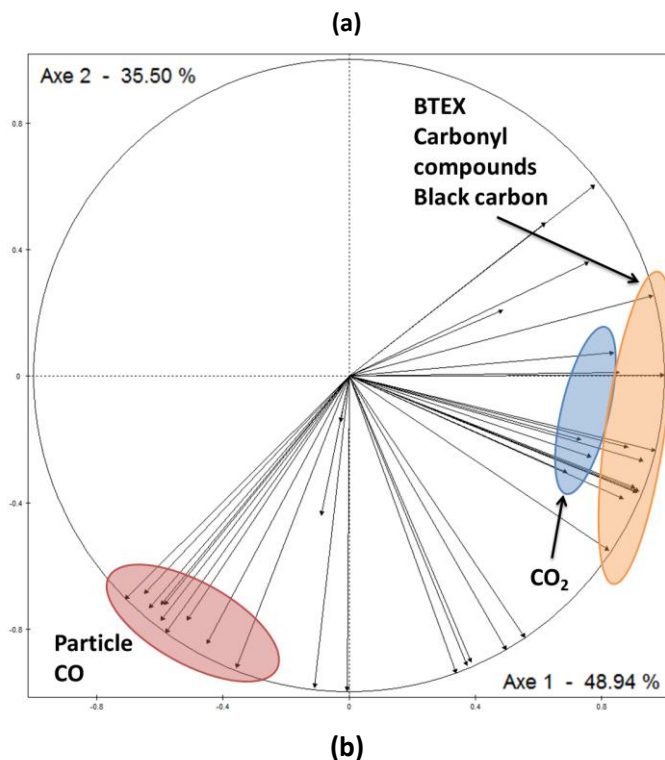
330 The impacts of driving conditions on pollutant emissions are studied using the PCA analysis with  
331 emission factors measured under the Artemis urban cold and hot start, Artemis road, Artemis  
332 motorway, and NEDC driving cycles. Two PCA analysis were performed, one for the Euro 4–6 diesel  
333 vehicles and one for the Euro 4–6 gasoline vehicle.

334 For the Euro 4–6 diesel vehicles, the urban cold start driving condition produces the most  
335 important emissions of various pollutants (PN, BTEX and carbonyl compounds...) compared to other  
336 driving conditions. For the Euro 4–6 gasoline vehicles, the impacts of the driving conditions on  
337 pollutant emissions are shown in the figure 5. Figure 5a shows the projection of the variables, the  
338 measured emission factors in this case, on the two principal axes. They account for 85% of the total  
339 variance, with a strong Axis 1 (49% of the variance). Figure 5b shows the projection of the driving  
340 cycles (individuals) in the same two dimensions.

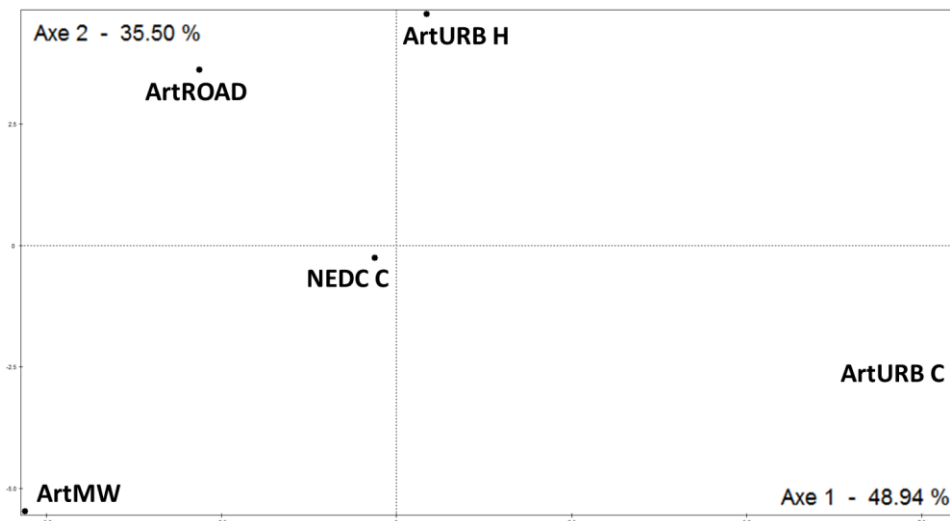
341 A cluster comprising BTEX, black carbon, and carbonyl compounds can be seen on the right  
342 portion of the correlation circle (Figure 5a). These compounds follow the same emission pattern and  
343 are at the same position as the Artemis urban driving cycle with cold start (Figure 5b). This  
344 observation indicates that the cold urban driving condition produces highest emissions of these  
345 compounds. Similar results were observed by Caplain et al. (26) and Louis et al. (14). Another cluster,

346 located in the upper left portion of the correlation circle and corresponding to the Artemis motorway  
 347 driving cycle, comprises particles, and CO. The motorway driving condition at high speed produces  
 348 highest particle, and CO emissions. Finally, the NEDC driving cycle is in the middle of the four Artemis  
 349 driving cycles of the factorial plan. This observation indicates that cycle emissions are not correlated  
 350 with the two principal axes.

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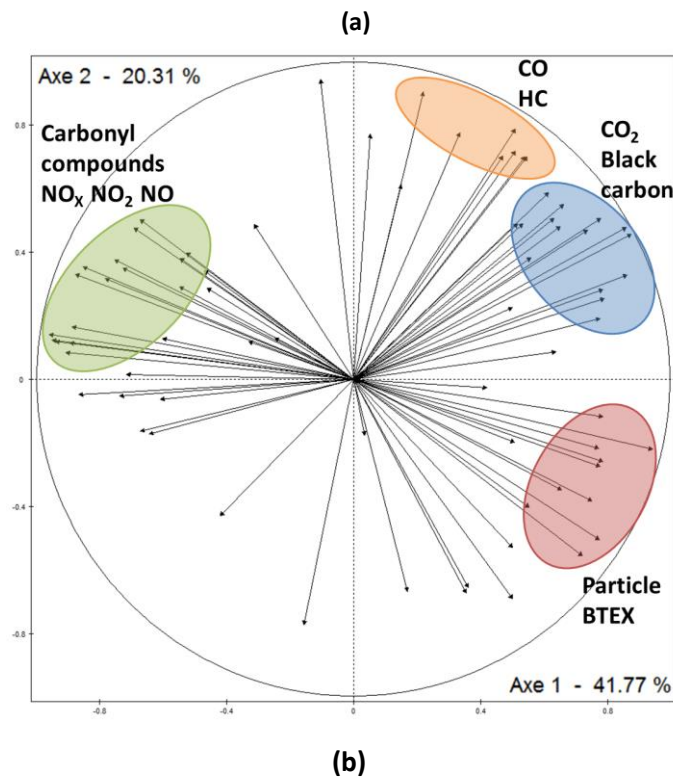
357 **Figure 5.** PCA performed for all the driving cycles for the Euro 4–6 gasoline vehicles, with (a) the  
 358 projection of emission factors and (b) the projection of cycles.

359 **2.4.2. Impacts of aftertreatment and engine technologies on pollutant emissions**

360 The eight vehicles tested were fitted with six different aftertreatment and engine technologies.  
 361 All three of gasoline vehicles in this study were fitted with TWC. However, they had different engine

362 technologies. The Euro 4 vehicle used an indirect injection technology, the Euro 5 vehicle used a  
 363 direct injection technology and the Euro 6 used a direct injection technology with propulsion engine  
 364 (located at the rear). All six diesel vehicles were fitted with DOC. Apart from this common system, the  
 365 aftertreatment technologies were different. One Euro 4 vehicle and one Euro 5 vehicle had a  
 366 catalyzed DPF, two Euro 5 vehicles had an additive DPF (add DPF), and one Euro 6 vehicle had a  
 367 catalyzed DPF (cat DPF) and NO<sub>x</sub> trap.

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374 **Figure 6.** PCA performed for all the driving cycles and all the vehicles, with (a) the projection of  
 375 emission factors and (b) the projection of vehicles.

376 To study the effects of vehicle technologies on the pollutant emissions, we performed PCA using  
 377 the emission factors measured for all the vehicles with all the Artemis and NEDC driving cycles as

378 variables, and the eight tested vehicles as individuals. Figure 6 shows the projection of the variables  
379 (Figure 6a) and individuals (Figure 6b) on the two principal axes. They account for 62% of the total  
380 variance, with a strong Axis 1 (42% of the variance). On the factorial plan, the diesel and gasoline  
381 vehicles are separated into two clusters. The gasoline vehicles are in the right portion of the plan and  
382 the diesel vehicles are in the left portion. In the case of the diesel vehicles, the two Euro 5 with  
383 additive DPF were clustered together and located at left bottom of the vehicle projection. In  
384 contrary, the three diesel vehicles with catalysed DPF located at top left of the plan. All three  
385 gasoline vehicles tested were located at different places: Euro 6 DI at right bottom; Euro 4 at right  
386 axis; and Euro 5 DI at top right.

387 The cluster on the left portion of the correlation circle comprises  $\text{NO}_x$  and carbonyl compounds.  
388 They correspond to the diesel vehicles fitted with a catalyzed DPF that meet Euro 4–6 standards  
389 which are characterized by higher emissions of these compounds. The Euro 6 Catalyzed DPF diesel  
390 vehicle with  $\text{NO}_x$  trap is located at the leftmost portion of the plan, indicating that this vehicle  
391 produces higher  $\text{NO}_x$  emissions. Such high emissions were also observed Ntziachristos et al. (35)  
392 under the WLTC cycle and PEMS measurements, between 100 and 1100 mg/km with high  
393 uncertainties. Two clusters at the upper right portion comprise CO and HC and black carbon and  $\text{CO}_2$ ,  
394 respectively. The cluster at the lower right portion comprises particles and BTEX. These various  
395 compounds are emitted in greater quantities by the Euro 4–6 gasoline vehicles compared to the  
396 diesel vehicles.

397 This preliminary PCA analysis shows the possibility of studying the impacts of driving conditions  
398 and technologies on emissions using a statistical method. However, the number of samples remains  
399 small and does not comprise all vehicle technologies. A detailed discussion is presented in the  
400 Section 4 herein.

## 401 **2.5. Discussion**

402 This paper aims to measure emissions of unregulated compounds from Euro 6 vehicles that are  
403 never measured in order to improve knowledge about their emissions under different driving  
404 conditions, and to supplement the first data for the emission factor inventory used by various  
405 emission models (HBEFA, COPERT IV). This paper also attempts to show whether it is possible to use  
406 PCA statistical analysis to investigate the impacts of aftertreatment technologies and driving  
407 conditions on vehicle emissions.

408 One Euro 6 diesel vehicle and one gasoline vehicle were tested on a chassis dynamometer for  
409 real-world Artemis, WLTC, and NEDC driving cycles. Emission factors of particles, black carbon,  $\text{NO}_2$ ,  
410 BTEX, PAH and carbonyl compounds for these vehicles were determined. The results show that diesel  
411 produces 200 to 5000 times more of  $\text{NO}_2$  emissions and 4 to 4500 times less of PN emissions than  
412 gasoline vehicle. Moreover, the Euro 6 diesel vehicle emits four times more carbonyl compounds  
413 than the Euro 6 gasoline vehicle. Compared to the Artemis urban hot start, Artemis urban cold start  
414 produces 100, 9–16, 2.5, and 10 times more PN, BTEX, carbonyl compounds, and BC emissions,  
415 respectively, and three times less  $\text{NO}_2$  emissions than Artemis urban hot start for both the diesel and  
416 gasoline vehicles. Compared to the WLTC approval cycle, diesel emission levels of PN, BC,  $\text{NO}_2$ , BTEX  
417 and carbonyl compounds are similar under the Artemis average cycles (urban cold start + road +  
418 motorway). For the gasoline vehicle, all the pollutant emissions are similar between the average  
419 Artemis and WLTC cycles. The only exception is BTEX: the WLTC cycle induced 4 to 13 times more  
420 emissions than average Artemis cycle. However, only two vehicles in this study were tested using the

421 WLTC cycle. Further testing will have to be conducted to confirm the impact of the WLTC driving  
422 condition on emissions of unregulated pollutants. Positive linear correlations have been observed for  
423 Euro 4–6 diesel vehicles between emissions of regulated compounds (HC and CO) and unregulated  
424 compounds (black carbon, benzene, toluene, xylene, formaldehyde and acetaldehyde). These  
425 correlations show that it is possible to estimate some of the unregulated compounds that are not  
426 always measured as a function of regulated compound emissions. These correlations only provide an  
427 approximate estimation that takes into account the large standard deviation.

428 PCA statistical analysis has been used as a preliminary method to see the possibility to  
429 investigate the impact of aftertreatment device and driving condition on pollutant emissions of eight  
430 vehicles with seven different technologies tested in this work and our previous work. With our PCA  
431 analysis, we show that the Euro 4–6 gasoline vehicles are characterized by higher particle number  
432 emission and lower NO<sub>x</sub> emission comparing to Diesel DPF vehicles. This statistical analysis with our  
433 data set is in agreement with Bach et al. (13), Alves et al. (23) and Louis et al. (15). They observed  
434 that Euro 4–5 gasoline vehicles emit around 100 times more particles and around 10 times less NO<sub>x</sub>  
435 comparing to the Euro 4–5 diesel equipped with DPF. The PCA analysis of the driving conditions  
436 showed that the urban driving conditions with cold start produce significant PN, BTEX, and carbonyl  
437 compound emissions compared to hot start for the Euro 4–6 diesel vehicles. This is confirmed by the  
438 results of Louis et al. (15), Caplain et al. (26) and Joumard et al. (21). The PCA showed that high-speed  
439 driving conditions (motorway) produce significant CO and PN emissions compared to low-speed  
440 driving conditions (urban) mainly induced by the gasoline vehicles. This is in agreement with Huang et  
441 al. (22). They observed that the urban driving condition with cold start emits around 2 to 150 times  
442 more BTEX and around two times more carbonyl compounds compared to the urban driving  
443 condition with hot start. For the impacts of aftertreatment and engine technologies, the PCA showed  
444 that diesel vehicles equipped with additive and catalyzed DPF emit few particles. Two different DPF  
445 technologies exhibit slightly different emission behaviors. The Euro 6 vehicle fitted with a NO<sub>x</sub> trap  
446 emitted the most NO<sub>x</sub>. The three gasoline vehicles, which are located opposite the diesel vehicles on  
447 the factorial plan, emit little NO<sub>x</sub>, but more particles, black carbon, BTEX, CO, HC, and CO<sub>2</sub>. Due to  
448 different engine technologies – indirect injection for Euro 4, direct injection for Euro 5 and direct  
449 injection with a propulsion engine for Euro 6 – the three gasoline vehicles exhibit different emission  
450 behaviors. These preliminary tests showed that different vehicle technologies or driving conditions  
451 can be characterized by some key pollutants. However, these tests included a relatively low sample  
452 number and did not cover all vehicle technologies. To complete our PCA analysis, we attempted to  
453 integrate results from other studies. For example: Caplain et al. (26) tested pre-Euro to Euro 3 diesel  
454 and gasoline vehicles. They monitored 11 aldehydes and 2 ketones as pollutants under urban with  
455 hot start and motorway driving conditions. Rehn (25) tested Euro 2 to Euro 4 diesel and gasoline  
456 vehicles by measuring HC, NO<sub>x</sub>, CO, PAHs in particulate phase, BTEX, aldehydes and acetone under  
457 Artemis urban with hot and cold start, and road driving conditions. Alves et al. (23) tested Euro 3 to  
458 Euro 5 diesel and gasoline vehicles. They monitored HC, CO, NO<sub>x</sub>, and BTEX under Artemis urban with  
459 hot and cold start, and road driving conditions. However, none of the pollutants monitored in these  
460 studies were measured for all Artemis driving conditions. Moreover, the particle number, black  
461 carbon, and several organic compound emission factors were not measured for Pre-Euro to Euro 3  
462 Diesel and gasoline vehicles. We therefore lack all the necessary data to complete our PCA data set  
463 for the same pollutants under all driving conditions or for all vehicles. The PCA analysis with  
464 integration of literature data shows that the missing data, which introduced significant bias to the



465 PCA analysis, should be included in a future study so as to extend this statistical method to a larger  
466 vehicle population.

## 467 **Acknowledgments**

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469 French Environment and Energy Management Agency (ADEME).

## 470 **Supporting information available**

471 Fuel composition for all tested vehicles (Table S1); Analytical methods for BTEX, carbonyl  
472 compounds, and PAH samples in the gas and particulate phases with quantification limit and  
473 uncertainty (Table S2); Emission factors for unregulated pollutant from Euro 6 diesel and gasoline  
474 vehicles (Table S3); Technical characteristics of the eight diesel and gasoline vehicles used for  
475 statistical analyses (Table S4). This material is available free of charge via the Internet at  
476 <http://pubs.acs.org>.

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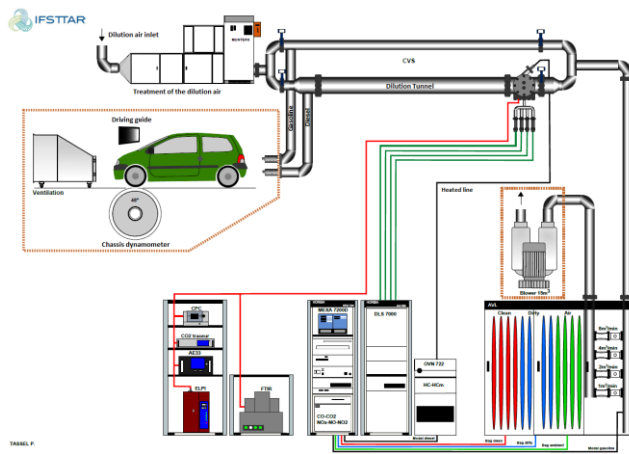
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