

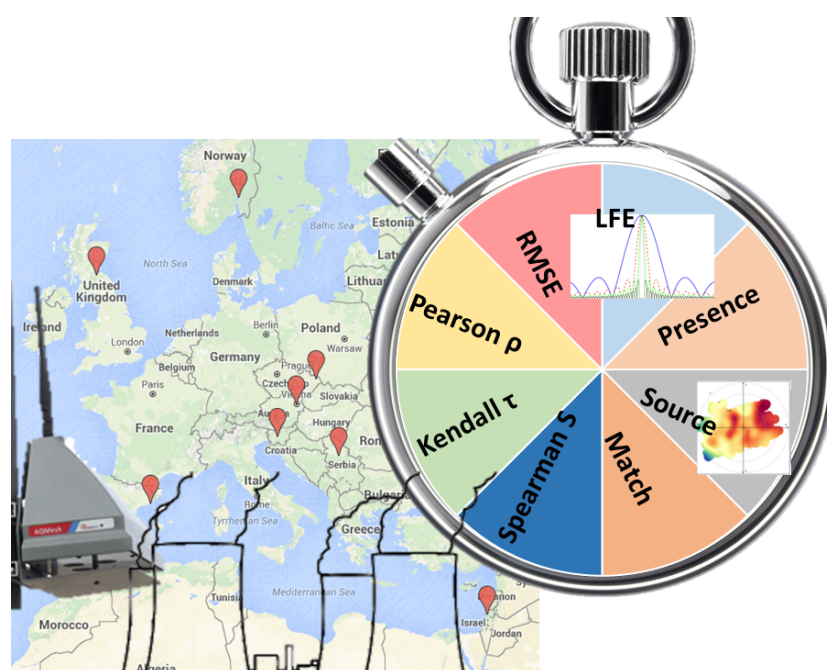
1 An Evaluation Tool Kit of Air Quality Micro-Sensing Units

2 Barak Fishbain^{1,*}, Uri Lerner^{1,†}, Nuria Castell-Balaguer², Tom Cole-Hunter^{3,4}, Olalekan Popoola⁵, David M.
3 Broday¹, Tania Martinez Iñiguez^{3,4}, Mark Nieuwenhuijsen³, Milena Jovasevic-Stojanovic⁶, Dusan
4 Topalovic^{7,6}, Roderic L. Jones⁵, Karen Galea⁸, Yael Etzion¹, Fadi Kizel¹, Yaela N. Golumbic^{9,1}, Ayelet
5 Baram-Tsabari⁹, Tamar Yacobi¹, Dana Drahlér¹, Johanna A. Robinson¹⁰, David Kocman¹⁰, Milena Horvat¹⁰,
6 Vlasta Svecova¹¹, Alexander Arpacı¹², Alena Bartonova²

7

8 † Equally contributed

9

10
11

TOC 1 – Graphical Table of Contents

¹ The Technion Center of Excellence in Exposure Science and Environmental Health (TCEEH), Faculty of Civil and Environmental Engineering, Technion – Israel Institute of Technology, Haifa, Israel

² Norwegian Institute for Air Research (NILU), Kjeller, Norway

³ ISGlobal, Barcelona Institute for Global Health, Barcelona, Spain

⁴ Centro de Investigación Biomédica en Red de Epidemiología y Salud Pública (CIBERESP), Madrid, Spain

⁵ Centre for Atmospheric Science, Department of Chemistry, University of Cambridge, Cambridge, England

⁶ VINČA Institute of Nuclear Sciences, University of Belgrade, Belgrade, Serbia

⁷ School of Electrical Engineering, University of Belgrade, Belgrade, Serbia

⁸ Centre for Human Exposure Science, Institute of Occupational Medicine (IOM), Edinburgh, Scotland

⁹ Faculty of Education in Science and Technology, Technion – Israel Institute of Technology, Haifa, Israel

¹⁰ Department of Environmental Sciences, Jožef Stefan Institute, Ljubljana, Slovenia

¹¹ Department of Genetic Ecotoxicology, Institute of Experimental Medicine AS CR, Prague, Czech Republic

¹² UBIMET GmbH, Vienna, Austria

12 Abstract

13 Recent developments in sensory and communication technologies have made the development of
14 portable air-quality (AQ) Micro-Sensing Units (MSUs) feasible. These MSUs allow AQ measurements in
15 many new applications, such as ambulatory exposure analyses and citizen science. Typically, the
16 performance of these devices is assessed using the mean error or correlation coefficients with respect to
17 a laboratory equipment. However, these criteria do not represent how such sensors perform outside of
18 laboratory conditions in large-scale field applications, and do not cover all aspects of possible differences
19 in performance between the sensor-based and standardized equipment, or changes in performance over
20 time. This paper presents a comprehensive Sensor Evaluation Toolbox (SET) for evaluating AQ MSUs by a
21 range of criteria, to better assess their performance in varied applications and environments. Within the
22 SET are included four new schemes for evaluating sensors' capability to: locate pollution sources;
23 represent the pollution level on a coarse scale; capture the high temporal variability of the observed
24 pollutant and their reliability. Each of the evaluation criteria allows for assessing sensors' performance in
25 a different way, together constituting a holistic evaluation of the suitability and usability of the sensors in
26 a wide range of applications. Application of the SET on measurements acquired by 25 MSUs deployed in
27 eight cities across Europe showed that the suggested schemes facilitates a comprehensive cross platform
28 analysis that can be used to determine and compare the sensors' performance. The SET was implemented
29 in R and the code is available on the first author's website.

30 1 Introduction

31 Air pollution is recognized as a contributing factor to various health outcomes, and has been associated
32 with public health risks [1, 2]. Accurately assessing ambient concentrations of different air pollutants is
33 necessary in any study on the impact of air quality (AQ) on different health endpoints. To date, ambient
34 pollutant concentrations are obtained from either short time-period measurement campaigns using a
35 large number of sensing devices (e.g. [3]), or from measurements reported by standard Air Quality
36 Monitoring (AQM) stations over extended time periods (e.g. [4]). While the former is limited in temporal
37 representativeness (e.g. due to inter-seasonal variation), the latter is limited in spatial representativeness
38 (e.g. due to dispersion patterns) and typically measures only a limited number of criteria pollutants [5].
39 Further, regulatory AQM stations require certified instrumentation meeting measurement accuracy
40 requirements, and an extensive set of procedures to ensure that data quality remains satisfactory. These
41 requirements, typically required by laws and regulations, ensure that measurements are comparable
42 across all networks with similar requirements, but limit the AQM spatial deployment due to their high

43 investment and operational cost. As a result, the AQM network has limited ability to account for spatial
44 variability of pollution levels in heterogeneous regions such as urban areas, which in return, renders
45 exposure assessment a very difficult task [6]. Moreover, the air-inlets of AQM stations are typically
46 located on rooftops or way above the ground [7], thus misrepresenting the true exposure of any individual
47 at head height.

48 Recent developments in sensory and communication technologies have made the deployment of portable
49 and relatively low-cost Micro Sensing Units (MSUs) possible. These MSUs can operate as a set of individual
50 nodes, or may be interconnected to form a Wireless Distributed Environmental Sensor Network (WDESN)
51 to measure air pollution over large spatial scales. WDESNs gather high-resolution spatial and temporal
52 data from numerous individual nodes allowing for a better interpolation and the generation of dense
53 pollution maps, which are closer to real-life pollution dispersion scenarios [8]. The gaseous sensors
54 mounted on these MSUs are low-power and low-cost, and are based on widely understood amperometric
55 sensor methodologies designed for sensing selected gases at the parts-per-million (ppm) level [9, 10, 11].
56 Electronic circuitry, which applies signal processing, allows for the detection at the part-per-billion level
57 [11]. Recent miniaturization of Optical Particles Counters (OPCs) [12, 13] and solid state [14, 15] sensors
58 allows to extend the MSUs capabilities to measure Particulate Matter (PM) as well.

59 The small size and low power-consumption of MSUs lay the path for many new applications that require
60 AQ data, such as exposure analyses [16, 17], education [18], hot-spot identification and characterization
61 [19], supplementary network monitoring [20, 21], and citizen science [22, 23, 24]. In particular, the
62 essence of citizen science requires active participation of citizens in the scientific research process [22].
63 Within the context of air-quality research, MSUs may be deployed at citizen's homes, monitoring either
64 ambient or indoor air quality in their local environment. An example is the CITI-SENSE project, which aims
65 at developing sensor- based citizen observatories for improving the quality of life in cities [25].

66 Seminal studies that evaluate MSUs in pre-field and field trials show that these units indeed can capture
67 air pollution spatio-temporal variation [26, 27, 11, 28, 21, 29, 30]. However, these studies have shown
68 that the MSUs' main limitation is their low accuracy relative to laboratory equipment [26, 27, 11, 28, 30]
69 or an AQM station [11, 28, 21].

70 Previously-used MSU calibration and evaluation measures, i.e., sensitivity [26, 27, 11, 28], correlation
71 coefficient, ρ , coefficient of determination, R^2 , [21, 11, 29], and the Root Mean Squared Error, RMSE [21,
72 29] aim at assessing the MSUs' accuracy and capability to capture trends and values of the pollutants' true
73 ambient levels. While these criteria evaluate some aspects of the sensors' performance in many fields, for
74 some applications different criteria covering additional performance aspects may be more adequate [24].

75 To date, some personal exposure studies have supplied participants with MSUs that measured various air
76 pollutants of exposure during daily routines (e.g., [2, 31]). However, exposure is affected by many factors,
77 and thus the variance of the dose response function is typically high and dominates the attributed relative
78 risks/hazard ratios results, regardless of sensors' accuracy [16, 17]. Therefore, one common practice for
79 estimating individual exposure is to use a coarse scale [5, 32, 25], rather than the sensors' actual
80 measurement. Educational and citizen science applications typically aim at fostering informal and
81 qualitative awareness. The measuring range in such applications is typically quantized into a binary scale,
82 indicating the presence or absence of a pollutant. These scales and measures, although quantized, can
83 still be used for relational comparison of air-pollution levels among different locations and times. This
84 motivates the need for a more widely composed set of criteria to characterize the MSUs' actual (field)
85 capabilities. Having such criteria allows for custom-made assessment of sensor's performance looking at
86 properties that are important according to the task and application in hand.
87 This work presents a comprehensive Sensor Evaluation Toolkit (SET) for evaluating and comparing the AQ-
88 MSUs' performance and its application on 25 sensors deployed in eight cities in Europe, as part of the
89 CITI-SENSE project [25]. The R implementation of the SET is available on the first author's website.

90 2 Material and Methods

91 MSU evaluation can be executed either in a laboratory, with critical atmospheric ambient conditions
92 measured and controlled, or in an open uncontrolled environment. The laboratory provides calibration
93 against traceable reference standards. In an open, uncontrolled environment, the MSUs are placed in
94 AQM stations and their measurements are compared against those acquired by AQM (reference)
95 equipment. While the SET requires a reference device (dubbed REF) to evaluate the MSU measurements,
96 it does not make any assertion on the nature of this reference equipment. The evaluation involves a
97 comparison of two concentration time-series: one acquired by the MSU, $\{C_k^{MSU}\}$, and one obtained by
98 the reference device, $\{C_k^{REF}\}$. Both time-series should be of equal length, i.e. consist of K measurements,
99 with the measurement acquired more or less simultaneously.

100 The SET consists of eight performance measures, including the classic measures of RMSE and various
101 correlations (described in Section 2.1). Four new measures, within the SET, are introduced: the *presence*
102 measure that represents the sensor's availability over time (described in Section 2.2); the *source-analysis*,
103 *which* depicts how accurately a sensor can identify and locate a source (detailed in Section 2.3); the *match*
104 (detailed in Section 2.4) that evaluates the sensor's accuracy when the measured concentrations are
105 transformed into generalized coarse scales; and the *Lower Frequencies Energy Content (LFE)*, which

106 measures the MSUs' ability to capture the temporal variability of the observed pollutant (Section 2.5). All
107 measures are then combined into an Integrated Performance Index (IPI) (Section 2.6).

108

109 2.1 Root Mean Squared Error and Correlation Coefficients

110 The Root Mean Squared Error (RMSE) and the Pearson correlation measures are often used to evaluate
111 MSUs' performance [8, 11, 21, 28, 29]. However, these measures apply specific assumptions on the errors
112 and their distributions. RMSE measures the total bias (deviation) between two time series, and is often
113 used to evaluate MSU errors [21, 28, 11]. While the RMSE is an excellent general-purpose error metric for
114 numerical deviations, it severely amplifies and disproportionately accounts for large errors. Thus, if two
115 signals have the same values but a small abrupt large deviation the inter-unit RMSE will be large.

116 Correlation coefficients are often used for evaluating the similarity between two time series, usually in
117 complement to the RMSE [21, 11, 28]. Correlation coefficients are robust to abrupt, large deviations and
118 are bounded between $[-1, 1]$, a property which will be exploited in our aggregation process (see
119 Section 2.6). Typically, the correlation of the tested device with a reference is reported with the Pearson
120 correlation coefficient (e.g., [8, 21, 11, 28, 29]). However, the commonly used Pearson correlation
121 coefficient, ρ , measures how well AQM measurements can be represented by the MSU records using a
122 linear function. This measure is adequate if both the AQM and the MSU are in their linear sensitivity range.
123 Ambient pollutant levels are often below the linear range of the MSUs [26, 27, 30, 28]. Therefore, the SET
124 includes also the Kendall- τ [33, 34] and the Spearman rank correlation coefficients, S , [35], which do not
125 assume normality of the underlying variables and perhaps more importantly, are more sensitive to
126 monotonic but non-linear relationships. A real-life example of using the differently defined correlation
127 coefficients is given in Section S1 of the supplementary information. The example presents seven NO
128 MSUs (CitiSense Leo Model, Ateknea Solutions Catalonia, Spain), which are evaluated against a reference
129 AQM for 20 days. While the Pearson and Kendall- τ coefficients are relatively low, and would have
130 rendered the sensor as inadequate, the Spearman coefficient shows that the sensor is suitable for NO
131 measurements, given linearity is not considered. This phenomenon is due to the low NO ambient levels
132 out of the linear response range of the MSUs [36]. Thus, the multiple evaluation criteria allow to better
133 characterize MSUs' suitability to different applications.

134

135

136 2.2 Presence

137 Each sensor's time series may contain missing values. This may result from the sensor malfunctioning or
138 from communication errors. The *presence* measure accounts for the sensor's or system's availability of a
139 measurement at a given time, and reports the fraction of the acquired measurements of all theoretically
140 possible. The presence of an MSUs is a significant factor in evaluating any outdoor measuring device. For
141 calculating, for example, averages during a given time interval, a minimum data availability is required to
142 ensure representativity. Limited presence always brings about the question of representativity of the
143 measurements for a given environment, and may indicate high maintenance costs. With that, sensor's
144 presence is completely omitted from the RMSE and correlation evaluation criteria. Standard AQM stations
145 are bounded to standards, connected to the power grid and are placed in dedicated containers, where
146 only the inlets are exposed to outside and weather conditions. Thus, measuring equipment presence is
147 typically a non-issue for AQMs. When it comes to MSUs, presence is often a major hurdle. This is why this
148 measure has a larger significance when evaluating MSUs.

149

150 2.3 Source Analysis

151 For many applications such as source apportionment [37, 38] or dispersion models, especially Lagrangian
152 models [39, 40], the source location is crucial. Bivariate polar plots, which represent how the
153 concentration of a pollutant varies with the wind direction and speed at the receptor, have proved to be
154 a useful tool for identifying and understanding pollution sources [37, 38, 39, 40, 21]. This representation
155 manifests the directional dependence of different sources, particularly when more than one monitoring
156 site is available, making *source-analysis* ideal for WDESN applications.

157 The source location analysis within the SET aims at evaluating how accurate the MSU is in identifying and
158 locating sources. i.e., it assesses the ability of the device to react to changes in observations within a time
159 interval that corresponds to wind direction change, and to be sensitive enough to measure associated
160 changes in concentrations. This is achieved through the calculation of the two-dimensional Pearson
161 correlation between polar plots obtained from the reference device and the MSU, treating both as two-
162 dimensional matrix arrays [41]. For generating the polar plots, time-matched measurements of the wind
163 and the pollutant must be available. This information is typically obtained either from the AQM station
164 (given it measures these meteorological parameters) or by an externally collocated wind vane. Section S2
165 of the Supplementary material presents an example of a set of two PM_{2.5} MSUs evaluated against an AQM
166 station. While all other performance criteria of the two sensors are relatively similar, the source analysis
167 score of the two sensors does show a difference. This may be attributed to the sensor being placed

195 incorrectly so it has an obscure observation. Thus, a low source analysis score may allow us to find the
196 problem and rectify it.

197

198 2.4 Match Score

199 Integrated AQ measures, such as the Air Quality Index (AQI) [32], are often used to convey the general
200 notion of severity of air-quality to the public, ranking observations according to a chosen scheme. Such
201 measures may also be used when the research question does not require precise measurements but
202 rather a more general interpretation, such as general risk estimation [5, 32] and citizen science [24]. When
203 applying such an AQ grading scheme, neither the RMSE nor the coefficient of determination represent
204 well the sensor's performance, as they penalize small perturbations in the measurements. Thus, if an MSU
205 deviates, due to its inaccuracy, from a reference device and if its deviations are randomly distributed
206 around the reference value, these two measures will report poor performance. The *match score*
207 overcomes this limitation and its calculation is as follows:

1. Set $COUNT = 0$
2. Compute the dynamic range for $\{c_t^{MSU}\}$ and for $\{c_t^{REF}\}$, i.e., $[\min\{c_t^{MSU}\}, \max\{c_t^{MSU}\}]$
and $[\min(\{c_t^{REF}\}), \max(\{c_t^{REF}\})]$ for all $t \in T$.
3. For $d = 1$ to D do:
 - a. Divide MSU's dynamic range into d equal bins and label them 1 through d .
Divide REF's dynamic range into d equal bins and label them 1 through d .
 - b. For each pair of measurements $\{c_t^{MSU}, c_t^{REF}\}, t \in T$:
If c_t^{MSU} and c_t^{REF} belong to the bins with the same label $\rightarrow COUNT = COUNT + 1$;
4. Compute: $Match\ Score = \frac{1}{D \cdot T} \cdot COUNT$

208

209 *Algorithm 1 – Match Score*

210 As can be seen, the match score is the proportion of agreement among strata for increasing amount of
211 sub-partitions between the reference and the MSU measurements. If we would like to compare the
212 different MSUs, the number of maximum sub-partitions, D , should be predetermined and be kept
213 constant throughout the analyses. Its value should be the highest number that still has at least one set of
214 measurements c_t^{MSU} and c_t^{REF} that belong to the same bin. In our analyses, after several preliminary
215 runs, D was set to 10.

216

217 **2.5 Lower Frequencies Energy (LFE)**

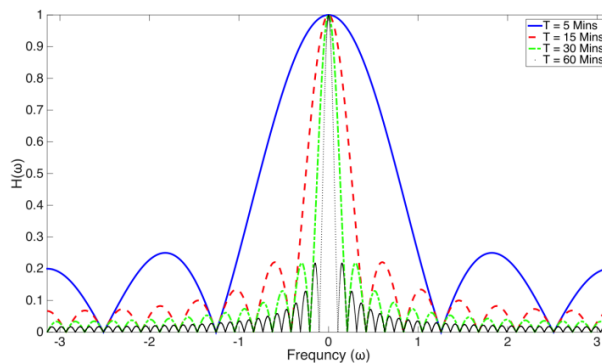
218 The signal Lower Frequency Energy, dubbed LFE, is a characteristic of the signal rather than a comparative
 219 measure with respect to a reference device. Let us assume that $p(t)$ is a continuous signal that represents
 220 the true ambient level of a specific pollutant in a specific location. Both the MSU and the AQM average
 221 $p(t)$ over a small sliding temporal-window, $h(t)$, of a size Δt , obtaining $\bar{p}(t)$, and sample it to obtain the
 222 aforementioned discrete time series $\{C_k^{MSU}\}$ and $\{C_k^{REF}\}$. Formally, the averaging is described by a
 223 convolution integral:

$$\bar{p}(t) = \int_{-\infty}^{\infty} p(\tau)h(t - \tau)d\tau \quad \text{Equation 1}$$

224 Let ω be the Fourier Transform domain coefficients. Applying the Fourier Transform on $p(t)$ and $h(t)$
 225 (obtaining $P(\omega)$ and $H(\omega)$) respectively and the convolution theorem [47], the Fourier representation of
 226 $\bar{p}(t)$, $\bar{P}(\omega)$, is given by:

$$\bar{P}(\omega) = P(\omega) \cdot H(\omega) \quad \text{Equation 2}$$

228 The Fourier Transform's amplitude of $h(t)$ is presented in Figure 1 for four different window sizes:
 229 averaging over 5, 15, 30 and 60 minutes. It can be seen that at zero $H(\omega)$ receives a value of one and its
 230 value decreases as ω (in absolute value) increases. Considering Equation 2, the sampling process
 231 suppresses higher frequencies, i.e, it applies low-pass filter on the observed signal. Thus, larger window-
 232 sizes in the signal domain, i.e. larger Δt , represent narrower filters in the frequency domain.



233
 234 *Figure 1 - Fourier Transform of $h(t)$ averaging over 5 min period, i.e. $\Delta t = 5$ (solid blue), $\Delta t = 15$ min (dashed red); $\Delta t = 30$ min (dot-*
 235 *dash green); and $\Delta t = 60$ min (dotted black).*

236 If the observed pollutant signal changes rapidly, its higher-frequency coefficients will assume higher
 237 values. Considering Equation 2, these values would be diminished in the acquired signal if they are

238 multiplied by a narrow $H(\omega)$, i.e. averaged over a large temporal-window. A real-life example is given in
239 Section S3 in the supplementary material.

240 The signal's energy is a characteristic used in signal processing and is given by:

$$E = \int_{-\infty}^{\infty} |p(t)|^2 dt \quad \text{Equation 3}$$

241 Following the Parseval's theorem [48], the energy of a signal is equal to the energy of its Fourier transform:

$$E = \int_{-\infty}^{\infty} |P(\omega)|^2 d\omega \quad \text{Equation 4}$$

242 Hence, the function $|P(\omega)|^2$ represents the energy distribution in the frequency domain. The smaller the
243 energy portion in the higher frequencies, the better the sensor can capture the signal's temporal
244 variability. Thus, the portion of the signal's energy in the lower frequencies, i.e. the *Lower Frequencies*
245 *Energy* (LFE), can be used for evaluating the sensor's capability to capture the temporal variability of the
246 pollutant. After discrete sampling that accounts for the K samples in the pollution time series Equation 4
247 becomes:

$$E = \sum_{\omega=1}^K |P(\omega)|^2, \quad \text{Equation 5}$$

248 and the LFE measure is computed as:

$$LFE = 1 - \frac{\sum_{\omega=1}^K (\omega \cdot |P(\omega)|^2)}{E \cdot \frac{K(K-1)}{2}} \quad \text{Equation 6}$$

249 where E is given by Equation 5.

250 The maximum value that LFE can obtain is 1, which represents the case where all the information is within
251 the first frequency coefficient, i.e., all frequency coefficients, but the first one, are zero.

252 MSUs are typically self-contained units with their own power supply and transmission modules. Typically,
253 data acquisition and transmission times are set such that operational energy consumption (for data
254 acquisition and transmission) is minimized. Consequentially, their sampling interval, Δt , may be long, i.e.,
255 low sampling rate, which corresponds to narrow low-pass filtering. Therefore, while applicable to AQM
256 and standard laboratory equipment, a measure of spectral distribution is especially important to MSUs
257 working under power consumption constraints.

258 2.6 Integrated Performance Index (IPI)

259 The SET consists of eight different performance measures accounting for different aspects of signal
260 acquisition. Different combinations of these measures can be used in order to evaluate the sensor
261 performance, depends on the specific application. In order to integrate the various measures, it is

262 important that they all share the same scale. This is inherent for the SET as all measures span between [0,
263 1].

264 Integration of several measures into an overall evaluation measure can be done either by addition or
265 multiplication of all measures together. The former facilitates an aggregation scheme that can account for
266 different weights for the different measures by introducing weight coefficients, $\vec{\alpha}$. Given $\vec{\alpha}$, and \vec{m} the
267 measures vector for a given sensor (RMSE, correlations, presence, source analysis, match and LFE) and
268 the time series $\{C_k^{MSU}\}$ and $\{C_k^{REF}\}$ acquired by the sensor and a reference device respectively, the IPI is
269 given by:

270

$$IPI^{MSU} = \sum_i \alpha_i m_i(\{C_k^{MSU}\}, \{C_k^{REF}\}) \quad \text{Equation 7}$$

271 2.7 SET Implementation and Application

272 For demonstrating the method and its capabilities, the SET was implemented in R. For evaluation, the
273 MSUs are compared against a reference device. This device can be either AQM or laboratory calibrated
274 equipment. Both the MSU and the reference device must measure the same physical phenomenon (e.g.,
275 ambient levels of a specific pollutant, temperature or relative humidity). The same physical phenomenon
276 can be measured when the sensors are collocated [21, 11, 28] or when the observed phenomenon is
277 uniform in all measuring points [21, 49]. When no AQM nor reference devices are available the same
278 analysis can be done with respect to the average signal of the entire sensory network in a given region
279 [21]. Here we demonstrate the SET for collocated sensors with AQM stations.

280 For demonstrating the capabilities and richness of the suggested evaluation toolkit, twenty-five MSU pods
281 (Geotech AQMesh, UK [50]) were placed near ten different AQM stations in eight cities in Europe, as part
282 of the European Union 7th framework program (FP7) CITI-SENSE project [25]. The full deployment,
283 acquiring data for about three months at each location, is detailed in Section S4 of the supplementary
284 material. Each AQMesh unit was equipped with five environmental sensors: NO, NO₂, O₃, atmospheric
285 pressure (AP), and relative humidity (RH). Some of the AQMesh pods included also OPC PM sensor.
286 Additionally, the AQMesh measured the unit's (internal) temperature (Temp). The specific AQM
287 parameters (location, height above ground level (AGL) and above sea level (ASL)) are detailed in Section
288 S4 of the supplementary information. The average temperature and the averages of all measured
289 pollutants are provided in Table 1 alongside their SET performance. The latter is color coded to represent
290 low to high SET values in a red-to-green color scale.

291 In order to compare the AQM and the MSU measurements, the time resolution of both should be the
292 same. If that is not the case, the finer time resolution time series has been aggregated so it fits the coarser
293 resolution. The MSU time-series were acquired at a 15-min resolution, while the AQM time-series had a
294 30 (or 60) -min resolution. Hence, MSU measurements were averaged (without overlap) to produce a
295 time-series that corresponds to the AQM temporal resolution.

296 3 Results and Discussion

297 3.1 Overview

298 Table 1 depicts the average values of the measured environmental parameters, showing that the MSUs'
299 meteorological measurements are more accurate than those of pollutant concentrations. The AP, (pod
300 internal) Temp and RH sensors have, on average, an Integrated Performance Index (IPI) of 0.975, 0.875
301 and 0.851, respectively. Among the pollutants, NO sensors had the highest IPI, with an average of 0.705.
302 O₃, CO and NO₂ obtained IPIs of 0.664, 0.609 and 0.578, respectively.

303 The utilization of the SET for evaluating MSU performance is well demonstrated in Table 1. Sensor 143,
304 which presents lower IPI values for all measured environmental parameters, may have experienced a
305 systematic error. This may result from incorrect placement of the sensor or malfunction of hardware.
306 Sensor GAP 4 presents low IPI for RH. The average RH value that this sensor reported was 106.4%. This
307 clearly suggests that the sensor is faulty or overly-offset for this parameter. Sensor 118 presented low IPI
308 for CO and NO while their average concentrations were much higher than those measured by the AQM
309 and other collocated MSUs. All these measurements were removed from the following analysis.

310 The richness offered by the IPI is presented in Table 2, through the breakdown of the IPI measure into its
311 components (Mean (M), Match score, RMSE, Pearson ρ , Kendall τ and Spearman (S) correlation
312 coefficients; Source-analysis score, Presence (Pres.) and Lower Frequencies Energy (LFE) content) for two
313 sensors – #118 NO and #130 Temp sensors. For both sensors the LFE measure is high, suggesting that the
314 changes in the observed signal are slower than the sampling rate. The #118 NO sensor presents extremely-
315 low correlation values, while its match score is high. Thus, while this specific sensor would grade poorly
316 using the traditional evaluation tools (correlation and RMSE), it would be more than sufficient for many
317 of the aforementioned applications, such as citizen science and exposure estimations. The Temp sensor
318 of pod #130 also presents interesting behavior. Its Match as well as its correlation coefficients are
319 reasonable, but its RMSE score is very low. This suggests that while the sensor does not represent the true
320 ambient levels, i.e., it has some bias, it does represent the signal's behavior (i.e., good correlations).
321 Indeed, this was the case, as explained in section 3.3. Therefore we conclude that the different

322 components of the IPI measure do give a better understanding of the sensor performance and its
323 suitability for different applications.

324 3.2 Temperature Impact on the Measurements

325 Ambient temperature has been pointed-out as a major factor affecting sensor performance [21, 11, 28,
326 29]. Here we examine this using the IPI. Figure 2 shows the average IPI of all 25 MSUs for all seven
327 measured parameters, as a function of the average temperature that was measured by the AQM station
328 throughout the campaign. The temperature effect is evaluated over 175 measurements. Each of these
329 175 measurements consists of more than three months' worth of data. Thus, the temperature evaluation
330 is based on a large dataset. No apparent trend is observed, suggesting that the MSUs manage to
331 compensate for any temperature impact on the measurements. Previously reported temperature effects
332 on AQ measurements may be attributed to higher pollution levels in winter time (due to higher pollution
333 source strength and pollution accumulation during periods with temperature inversions [51, 52]). This is
334 because while temperature was not found to affect sensors' performance, the measured ambient levels,
335 as is shown later, do have an effect, where the sensors performs better in higher pollution levels. Next,
336 we analyze the IPI specifically for each measured parameter.

337

Table 1 – Environmentally sensed indicators - mean values (M) and Integrated Performance Index (IPI) for Air Pressure (AP); Temperature (Temp); Relative Humidity (RH); nitrogen oxide (NO); nitrogen dioxide (NO₂); ozone (O₃); and carbon monoxide (CO)

Unit	Location	AQM Temp [°C]	AP		MSU Temp		RH		NO		NO ₂		O ₃		CO	
			M	IPI	M	IPI	M	IPI	M	IPI	Mean	IPI	M	IPI	M	IPI
GAP1	Gal·la	23.62	1007	0.949	24.5	0.887	61.6	0.863	6.2	0.656	13.6	0.464	133.8	0.687	162.9	0.587
GAP2	Plac·dia	23.62	1008	0.952	24.5	0.892	62.0	0.864	0.1	0.412	0.8	0.455	178.6	0.669	177.0	0.572
GAP3	Barcelona	23.62	1008	0.950	24.5	0.894	60.8	0.853	1.0	0.557	9.4	0.497	65.4	0.699	137.9	0.584
GAP4	Spain	23.62	1008	0.944	24.3	0.897	106.4	0.382	16.2	0.667	9.6	0.503	144.8	0.675	159.0	0.569
GAP5		23.62	1009	0.957	24.5	0.901	68.7	0.543	1.9	0.638	1.8	0.481	153.8	0.674	148.3	0.576
116	St Leonards,															
118	Edinburgh,	-- ¹³	-- ¹³	-- ¹³	-- ¹³	-- ¹³	-- ¹³	-- ¹³	7.8	0.513	11.7	0.409	51.2	0.649	58.8	0.524
120	Scotland								138.0	0.501	8.5	0.394	66.9	0.604	433.3	0.451
									9.7	0.514	9.9	0.413	42.5	0.627	54.1	0.482
135	Neve	12.95	-- ¹³	-- ¹³	14.7	0.717	57.5	0.860	3.2	0.513	2.7	0.621	71.2	0.663	103.2	0.562
136	Shaannan,	12.95	-- ¹³	-- ¹³	14.4	0.689	57.0	0.842	3.3	0.508	5.0	0.640	44.8	0.697	94.4	0.552
130	Haifa, Israel	12.97	-- ¹³	-- ¹³	13.6	0.712	57.3	0.829	7.2	0.620	-- ¹³	-- ¹³	52.3	0.641	-- ¹³	-- ¹³
134	Igud,	13.65	-- ¹³	-- ¹³	14.9	0.705	58.1	0.802	5.9	0.633	-- ¹³	-- ¹³	41.9	0.633	-- ¹³	-- ¹³
	Haifa, Israel															
125	Ljubljana,	12.46	979	0.989	13.7	0.935	64.9	0.947	-- ¹³	-- ¹³	21	0.55	78.4	0.711	166.4	0.769
128	Slovenia	12.46	978	0.940	13.8	0.926	64.8	0.945	-- ¹³	-- ¹³	3800	0.46	105.8	0.687	175.0	0.681
131		12.46	980	0.934	14.1	0.908	63.6	0.937	-- ¹³	-- ¹³	6	0.54	96.9	0.719	179.8	0.663
124		6.80	1009	0.988	7.9	0.931	84.2	0.889	27.2	0.921	15.5	0.685	-- ¹³	-- ¹³	101.2	0.710
144		6.80	1007	0.990	7.9	0.945	83.1	0.895	30.5	0.869	16.9	0.637	-- ¹³	-- ¹³	95.6	0.697
145	Kirkeveien,	6.80	1008	0.989	7.8	0.937	84.5	0.893	23.0	0.860	16.5	0.712	-- ¹³	-- ¹³	98.8	0.704
146	Oslo,	6.80	1008	0.988	7.9	0.937	84.2	0.896	37.3	0.899	13.7	0.697	-- ¹³	-- ¹³	102.1	0.682
147	Norway	6.80	1008	0.989	7.9	0.945	83.3	0.892	25.9	0.876	14.4	0.565	-- ¹³	-- ¹³	94.5	0.592
124		7.26	1006	0.976	9.1	0.923	65.9	0.923	16.8	0.859	9.2	0.583	-- ¹³	-- ¹³	-- ¹³	-- ¹³
144		7.26	1004	0.993	8.8	0.933	66.9	0.931	16.8	0.871	12.2	0.651	-- ¹³	-- ¹³	-- ¹³	-- ¹³

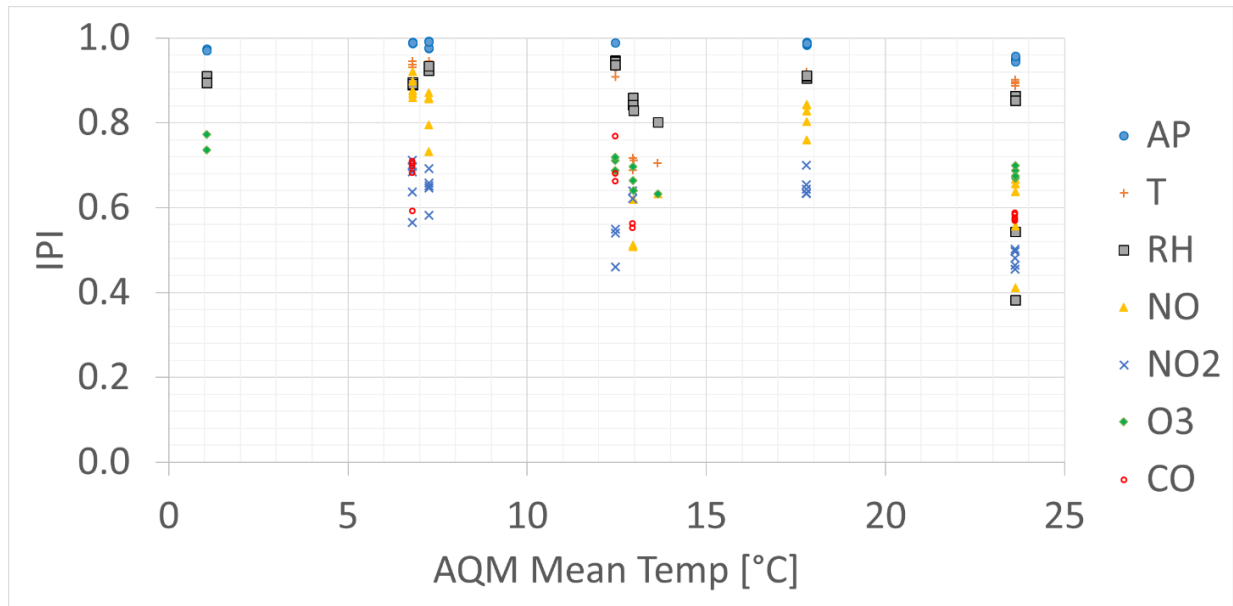
¹³ This specific environmental variable was not measured by the AQM at the collocation time period.

145	Hjortnes, Oslo, Norway	7.26	1004	0.992	8.7	0.937	67.0	0.930	11.9	0.795	16.5	0.692	-- ₁₃	-- ₁₃	-- ₁₃	-- ₁₃	
146		7.26	1005	0.978	9.1	0.924	66.2	0.923	9.8	0.732	12.6	0.647	-- ₁₃	-- ₁₃	-- ₁₃	-- ₁₃	
147		7.26	1005	0.991	8.8	0.945	66.4	0.934	16.4	0.856	9.3	0.659	-- ₁₃	-- ₁₃	-- ₁₃	-- ₁₃	
124		17.80	1011	0.983	20.4	0.911	63.0	0.907	15.4	0.842	17.7	0.654	-- ₁₃	-- ₁₃	-- ₁₃	-- ₁₃	
144		17.80	1009	0.991	20.5	0.905	62.8	0.905	15.5	0.843	22.2	0.644	-- ₁₃	-- ₁₃	-- ₁₃	-- ₁₃	
145		17.80	1010	0.989	20.4	0.918	63.7	0.911	5.9	0.760	26.9	0.700	-- ₁₃	-- ₁₃	-- ₁₃	-- ₁₃	
146		17.80	1010	0.986	20.5	0.917	63.1	0.910	22.5	0.803	18.7	0.635	-- ₁₃	-- ₁₃	-- ₁₃	-- ₁₃	
147		17.80	1010	0.988	20.6	0.920	61.9	0.912	12.7	0.828	15.6	0.634	-- ₁₃	-- ₁₃	-- ₁₃	-- ₁₃	
611		Ostrava, Czech Rep.	1.06	992	0.975	4.6	0.901	80.5	0.911	-- ₁₃	-- ₁₃	-- ₁₃	-- ₁₃	30.5	0.736	-- ₁₃	-- ₁₃
612			1.06	991	0.971	4.6	0.896	80.8	0.895	-- ₁₃	-- ₁₃	-- ₁₃	-- ₁₃	26.3	0.773	-- ₁₃	-- ₁₃
221	Belgrade, Serbia	5.78	1008	0.685	5.7	0.885	82.8	0.810	69.7	0.794	27.5	0.591	35.8	0.585	519.8	0.691	
222		5.78	1008	0.685	5.7	0.874	81.4	0.815	81.8	0.793	14.3	0.554	60.9	0.537	537.6	0.692	
143	Vienna	5.08			15.1	0.473	62.4	0.402	13.2	0.387	5.5	0.348	126.0	0.402	-- ₁₃	-- ₁₃	
Average IPI				0.953		0.876		0.848		0.711		0.577		0.653		0.617	

341 Table 2 – IPI breakdown – Mean ambient level (M); Match score; Root Mean Squared Error (RMSE); Pearson ρ correlation
 342 coefficient; Kendall τ correlation coefficient; Spearman (S) correlation coefficient; Source analysis; Presence (Pres.); Low
 343 Frequencies Energy (LFE) content and the integrated Air Quality Index (IPI)

Sensor	M	Match	RMSE	ρ	τ	S	Source	Pres.	LFE	IPI
118 (NO)	129.9	0.920	0.24	0.063	0.068	0.090	-- ¹⁴	0.732	0.976	0.519
130 (T)	13.63	0.462	0.003	0.679	0.538	0.712	-- ¹⁴	1	0.997	0.712

344



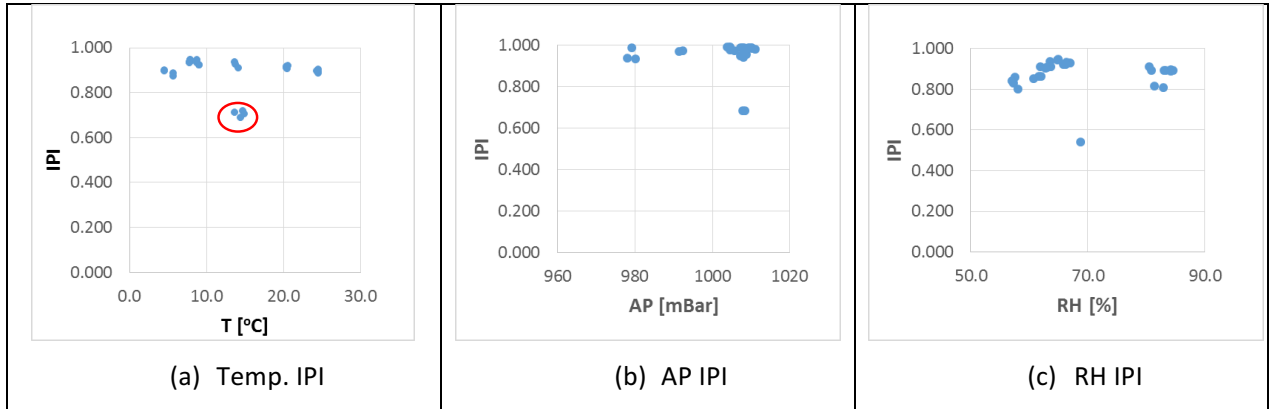
345

346 Figure 2 - IPI of the various measured parameters as a function of the ambient Temperature

347 3.3 Meteorological Sensors

348 Figure 3 presents the IPI for the Temp, AP and RH sensors for all MSUs, as a function of their average
 349 values. The first notion that arises from this presentation is that, in general, the sensors' IPI is indifferent
 350 to changes in the observed meteorological parameter. It is worthwhile mentioning that the batch of
 351 temperature sensors with lower IPI values (marked in circle in Figure 3a) were all obtained in the same
 352 station in Haifa, Israel. Further queries revealed that temperature records in this specific AQM were
 353 measured inside the monitoring station rather than outside, where the MSUs were located. This example
 354 demonstrates once again the use of the SET to point out sensing errors.

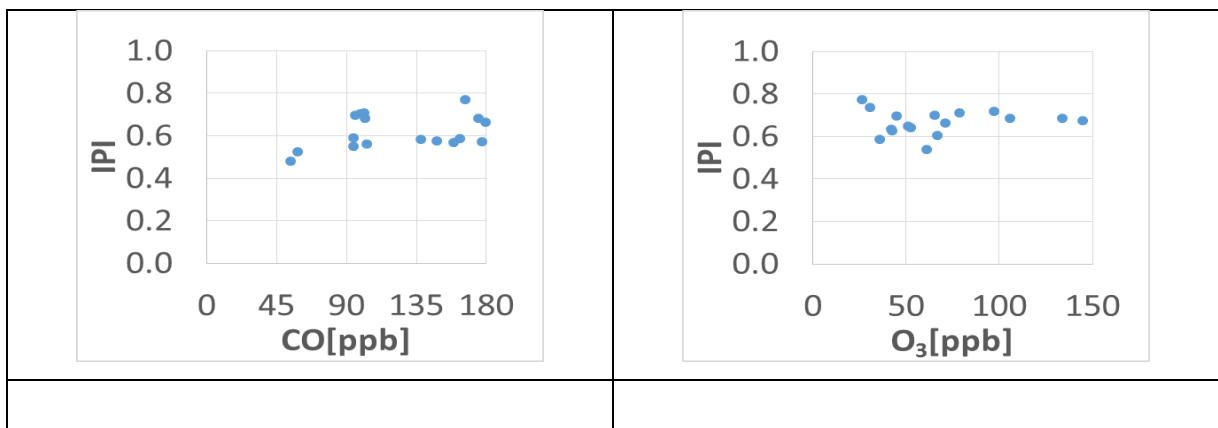
¹⁴ The Source score could not be computed as no wind data was collected at this location.

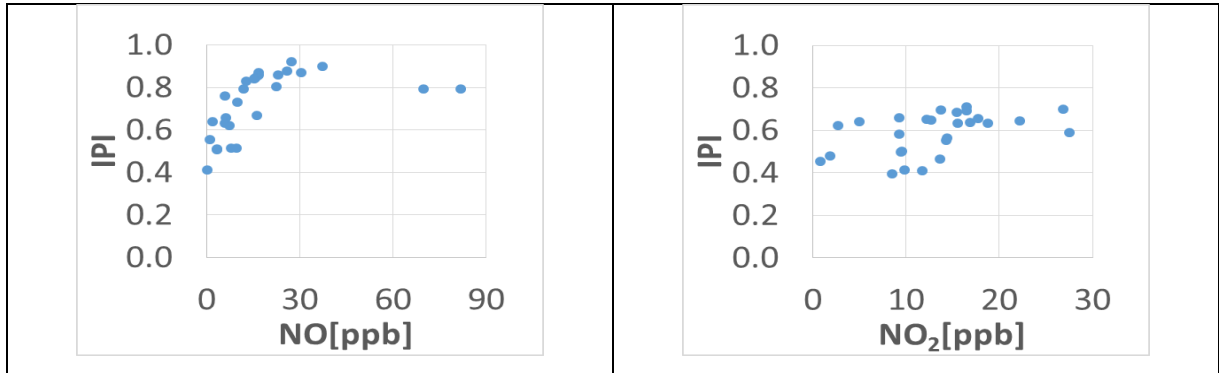


355 Figure 3 – IPI of a sensor as a function of its specific measured Meteorological Parameters – Temperature IPI (a), Air Pressure (AP)
 356 IPI (b), and Relative Humidity (RH) IPI (c)

357 3.4 Gaseous Pollutants Sensors

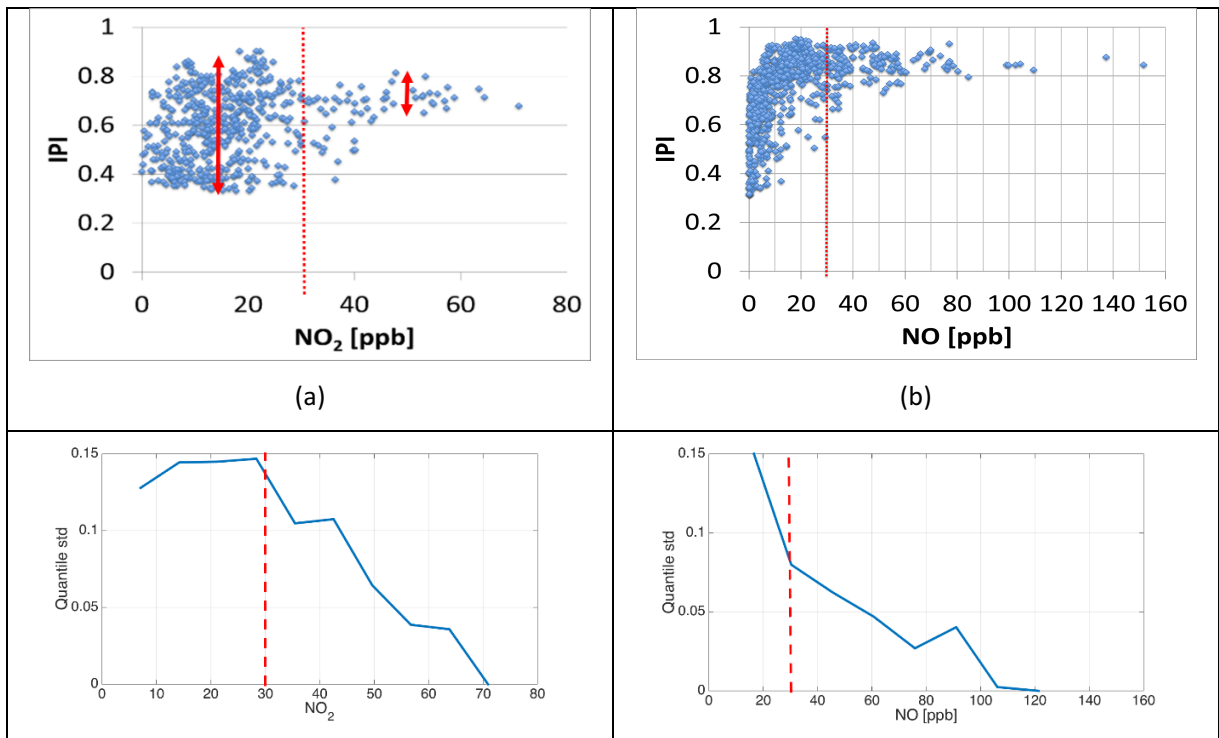
358 Figure 4 depicts pollutant specific sensor's IPI as a function of the average reading. Thus, the IPI of each
 359 pollutant's sensor, computed over the entire campaign, is plotted against the same pollutant's campaign
 360 average reading. It is evident from Figure 4 that the ambient level of the observed pollutant has a direct
 361 impact on the sensing quality. The lower the ambient pollutant level, the lower the IPI and the higher its
 362 variance for similar pollutant's ambient levels (i.e., the sensors presents lower reliability for lower
 363 pollutants ambient levels). Similar behavior was observed by Lerner et. al [29] and Moltchanov et. al [21].
 364 Hence, the IPI suggests that the MSU sensors are more suitable for locations where the pollutant is known
 365 to be high. Means to extract the threshold are discussed next. As different pollutants have different
 366 ambient levels, it is important to note that the x-axes of all following figures present different scales.





367 *Figure 4 -IPI as a function of Pollutants' Recorded campaign average levels for each sensor on each node*

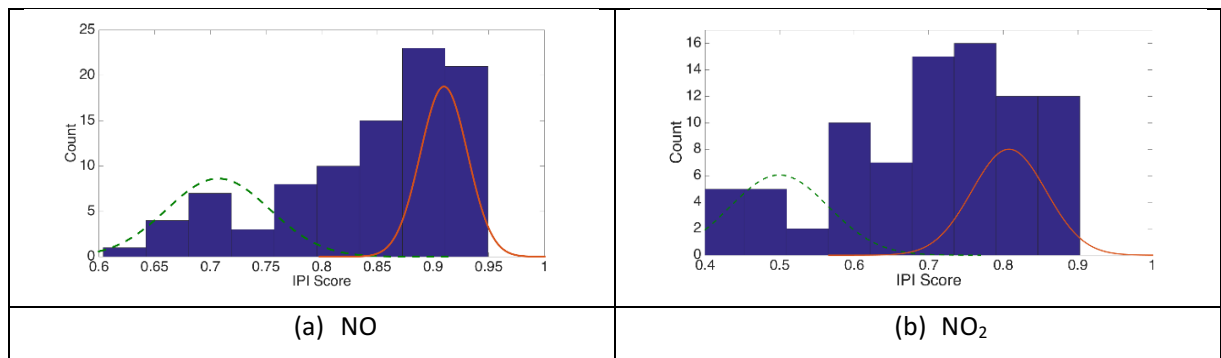
368 The aforementioned IPI behavior as a function of the pollutant's ambient level is better observed when
 369 the IPI is computed based on a daily time series rather than using the entire campaign's time series. For
 370 example, Figure 5, depicts the daily IPI for NO₂ (a) and NO (b) from measurements obtained in Kirkeveien,
 371 Oslo, by sensors #124, #144, #145, #146 and #147 (see Section S4). One should determine the minimum
 372 ambient levels a sensor can measure where the IPI measure is high and the standard deviation is low (how
 373 high and how low is application dependent). Figures (c) and (d) depict the standard deviation of the IPI,
 374 computed for the associated IPIs of each decile of the pollution levels. It is evident that the standard
 375 deviation decreases as the IPI increases. Using the notions above, a sensible threshold for the sensors
 376 described for measuring NO₂ and NO can be ambient levels that are higher than 30 ppb.





377 Figure 5 - IPI as a function of NO₂ (a) and NO (b) Recorded Levels at Kirkeveien, Oslo (daily score). The standard deviation of the
 378 IPI computed for each pollution level decile are presented in Figures (c) and (d) for NO₂ and NO respectively

379 Figure 6 demonstrates the utilization of the IPI for comparison between sensors and between different
 380 working conditions; The figure depicts the histograms of the NO and NO₂ levels at Kirkeveien, Oslo, which
 381 are presented in Figure 5 (as time series). The notion above, of the effect of ambient levels on the sensors'
 382 performance, is evident in Figure 6, where a bimodal distribution of the IPI is well observed. The two
 383 models correspond to IPI scores above and below 30 ppb. Having the bimodal distributions' parameters
 384 inferred, one can study their relations. Formally, this can be done by the Kolmogorov-Smirnoff test for
 385 discrete variables [53, 54] comparing the best fitting distribution with the empirical distribution and test
 386 for significance.

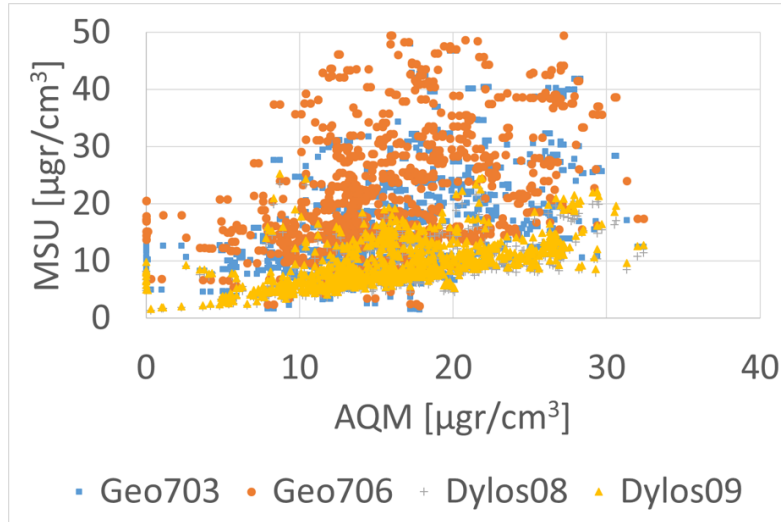


387 Figure 6 – NO (a) and NO₂ (b) IPI distribution functions measured at Kirkeveien, Oslo (daily score)

388 3.5 Particulate Matter Sensors

389 The SET criteria was applied to four different PM sensors, all collocated at the Igud AQM station (Haifa,
 390 Israel – see Table 1). The four sensors, two DC1700 Dylos (US) and two optical counters integrated on the
 391 GeoTech MSUs, were placed next to the AQM's inlet between December 17th and 24th, 2015. The data was
 392 recorded in 5 minutes' intervals. The four sensors measurements are displayed against AQM PM
 393 measurements (all measurements are in [$\mu\text{gr}/\text{m}^3$]).

394



395
396
397

Figure 7 – PM MSU measurements obtained between Dec. 17th and 24th, 2015 plotted against collocated AQM station.

398 Table 3 shows the IPI breakdown for the PM sensors. The two types of sensors present different
399 characteristics, while the GeoTech sensors present better RMSE, the Dylos ones have better match,
400 correlations and eBalance. Thus, the SET is also capable to evaluate PM sensors and define better the
401 suitability of a sensor to a specific application.

402
403

Table 3 – IPI breakdown of PM sensors

	Mean	Match	RMSE	Pearson	Kendall	Spearman	Presence	eBalance	IPI
Geo703	23.766	0.101	0.038	0.147	0.309	0.453	1.000	0.982	0.565
Geo706	26.926	0.119	0.023	0.259	0.284	0.421	1.000	0.993	0.579
Dylos08	139.894	0.311	0.134	0.560	0.468	0.639	1.000	0.997	0.693
Dylos09	152.258	0.312	0.147	0.570	0.474	0.647	1.000	0.997	0.695

404
405

406 4 Conclusions

407 This paper presents a Sensor Evaluation Toolbox (SET) for evaluating AQ MSUs by a range of criteria.
408 The rich evaluation provided by the suggested scheme allows for better assessment of sensors'
409 performance in varied applications and environments. The SET consists of eight different assessment
410 criteria: Root Mean Squared Error (RMSE), Pearson, Kandel and Spearman correlations, and four new
411 performance measures for evaluating sensors' capability to: locate pollution sources; represent the
412 pollution level on a coarse scale; capture the high temporal variability of the observed pollutant and their
413 reliability.

414 Application of the SET on measurements acquired by gaseous and PM MSUs deployed in eight cities across
415 Europe showed that each of the eight measures provides an important and unique information on the
416 sensor's performance assessing a rich spectrum of MSU capabilities. The result also demonstrate how the
417 scheme can pinpoint systematic as well as sensor's specific faults. Further, we demonstrated that the
418 Integrated Performance Index (IPI) can support a methodology for determining the sensors' performance,
419 hence facilitating a true cross platform evaluation. The SET was implemented in R [55] and is available on
420 the first author's website¹⁵.

421 Acknowledgements

422 We would like to acknowledge the AQM station managers for providing ambient air quality data, including
423 the Haifa District Municipal Association for Environmental Protection (Israel); Xarxa de Vigilància i Previsió
424 de la Contaminació Atmosfèrica de Catalunya (XVPCA), Direcció General de la Qualitat Ambiental,
425 Departament de Territori i Sostenibilitat, Generalitat de Catalunya (Spain); the Slovenian Environment
426 Agency (Slovenia); the Czech Hydrometeorological Institute (Czech Republic); Bureau Veritas, UK Ltd.
427 (Edinburgh); the Serbian Environmental Protection Agency – SEPA (Serbia).
428 CITI-SENSE, initiated in October 2012, is a four year Collaborative Project partly funded by the EU FP7-
429 ENV-2012 under grant agreement 308524.

430 References

431

- [1] International Agency for Research on Cancer (IARC), "Ambient Air Pollution," in *IARC Meeting Monographs*, Lyon, France, 2013.
- [2] J. Sarnat, P. Koutrakis and H. Suh, "Assessing the Relationship between Personal Particulate and Gaseous Exposures of Senior Citizens Living in Baltimore, MD," *Journal of the Air & Waste Management Association*, vol. 50, no. 7, pp. 1184-1198, 2000.
- [3] D. Crouse, M. Goldberg and N. Ross, "A prediction-based approach to modelling temporal and spatial variability of traffic-related air pollution in Montreal, Canada," *Atmos Environ*, vol. 43, no. 32, p. 5075–5084, 2009.

¹⁵ <http://fishbain.net.technion.ac.il> – will be available upon acceptance for publication

- [4] C. Pope, R. Burnett, M. Thun, E. Calle, D. Krewski, K. Ito and G. Thurston, "Lung Cancer, Cardiopulmonary Mortality, and Long-term Exposure to Fine Particulate Air Pollution," *JAMA*, vol. 287, no. 9, p. 1132–1141, 2002.
- [5] B. Bishoi and A. Prakash, "A Comparative Study of Air Quality Index Based on Factor Analysis and US-EPA Methods for an Urban Environment," *Aerosol and Air Quality Research*, vol. 9, no. 1, pp. 1-17, 2009.
- [6] S. Rao, V. Chirkov, F. Dentener, R. Dingenen, S. Pachauri, P. Purohit, M. Amann, C. Heyes, P. Kinney, P. K. Z. Kolp, K. Riahi and W. Schoepp, "Environmental Modeling and Methods for Estimation of the Global Health Impacts of Air Pollution," *Environmental Modeling & Assessment*, vol. 17, no. 6, pp. 613-622, 2012.
- [7] European Environment Agency, "Assessment and Management of Urban Air Quality in Europe," Office for Official Publications of the European Communities, Denmark, 1998.
- [8] P. Kanaroglou, M. Jerrett, J. Morrison, M. Bernardo Beckerman, A. Arain, N. Gilbert and J. Brook, "Establishing an air pollution monitoring network for intra-urban population exposure assessment: a location-allocation approach," *Atmos. Environ.*, vol. 39, no. 13, p. 2399–2409, 2005.
- [9] J. R. Stetter and J. Li, "Amperometric gas sensors a review," *Chemical reviews*, vol. 108, no. 2, pp. 352-366, 2008.
- [10] A. Bard and L. Faulkner, *Electrochemical Method: Fundamentals and Application*, 2 ed., New-York, NY, USA: John Wiley & Sons, 2001.
- [11] M. Mead, O. Popoola, G. Stewart, P. Landshoff, M. Calleja, M. Hayes, J. Baldovi, M. McLeod, T. Hodgson, J. Dicks, A. Lewis, J. Cohen, R. Baron, J. Saffell and R. Jones, "The use of electrochemical sensors for monitoring urban air quality in low-cost, high-density networks," *Atmospheric Environment*, vol. 70, pp. 186-203, 2013.
- [12] J. Ulanowski, P. Kaye, E. Hirst, A. Wieser and W. Stanley, "Miniature, low-cost optical particle counters.," in *Int. Conf. Optical Characterization of Atmospheric Aerosols*, Smolenice, Slovakia, 2013.
- [13] R. S. Gao, H. Telg, R. J. McLaughlin, S. J. Ciciora, L. A. Watts, M. S. Richardson, J. P. Schwarz, A. E. Perring, T. D. Thornberry, A. W. Rollins, M. Z. Markovic, T. S. Bates, J. E. Johnson and D. W. Fahey, "A light-weight, high-sensitivity particle spectrometer for PM_{2.5} aerosol measurements," *Aerosol Science and Technology*, vol. 50, no. 1, pp. 88-99, 2016.

- [14] M. Carminati, L. Pedalàa, E. Bianchi, F. Nasonb, G. Dubini and L. Cortelezzi, "Capacitive detection of micrometric airborne particulate matter for solid-state personal air quality monitors," *Sensors and Actuators A: Physical*, vol. 219, pp. 80-87, 2014.
- [15] M. Carminati, G. Ferrari and M. Sampietro, "Emerging miniaturized technologies for airborne particulate matter pervasive monitoring," *Measurement*, 2015.
- [16] E. Lebret, "Error in exposure measures," *Toxicol Indust Health*, vol. 6, p. 147–156, 1990.
- [17] M. Jerret, A. Arain, P. Kanaroglou, B. Beckerman, D. Potoglou, T. Sahsuvaroglu, J. Morrison and C. Giovis, "A review and evaluation of intraurban air pollution exposure models," *Journal of Exposure Analysis and Environmental Epidemiology*, vol. 15, pp. 185-204, 2005.
- [18] R. Ballantyne, J. Fien and J. Packer, "Program Effectiveness in Facilitating Intergenerational Influence in Environmental Education: Lessons From the Field," *The Journal of Environmental Education*, vol. 32, no. 4, pp. 8-15, 2010.
- [19] Y. Ma, M. Richards, M. Ghanem, Y. Guo and J. Hassard, "Air Pollution Monitoring and Mining Based on Sensor Grid in London," *Sensors*, vol. 8, no. 6, pp. 3601-3623, 2008.
- [20] The European Parliament and the Council of the European Union, "Directive 2008/50/EC on ambient air quality and cleaner air for Europe," *Official Journal of the European Union*, pp. L152/1-L152/44, 21 May 2008.
- [21] S. Molchanov, I. Levy, Y. Etzion, U. Lerner, D. Broday and B. Fishbain, "On Distributed Air Quality Measurements with Micro Sensors," *Science of the Total Environment*, vol. 502, pp. 537-547, 2015.
- [22] R. Bonney, H. Ballard, R. Jordan, E. McCallie, T. Phillips, J. Shirk and C. C. .. Wilderman, "Public Participation in Scientific Research: Defining the Field and Assessing Its Potential for Informal Science Education," 2009.
- [23] J. Shirk, H. Ballard, C. Wilderman, T. Phillips, A. Wiggins, R. Jordan, E. McCallie, B. Lewenstein, M. Krasny and R. Bonney, "Public Participation in Scientific Research : a Framework for Deliberate Design," *Ecology and Society*, vol. 17, no. 2, 2012.
- [24] R. Williams, V. Kilaru, E. Snyder, A. Kaufman, T. Dye, A. Rutter, A. Russell and H. Hafner, "Air Sensor Guidebook," National Exposure Research Laboratory, Office of Research and Development, U.S. Environmental Protection Agency, Research Triangle Park, NC, USA, 2014.
- [25] CITI-SENSE Project, "Home Page," 2015. [Online]. Available: <http://www.citi-sense.eu>. [Accessed 01 04 2015].

- [26] D. Lee and D. Lee, "Environmental gas sensors 1 (3), 214–224.," *IEEE Sensors Journal*, vol. 1, no. 3, pp. 214-224, 2001.
- [27] T. Becker, S. Muhlberger, C. Braunmuhl, G. Muller, T. Ziemann and K. Hechtenberg, "Air pollution monitoring using tin-oxide-based micro-reactor system," *Sensors Actuators B*, vol. 69, p. 108–119, 2000.
- [28] D. Williams, G. Henshaw, M. Bart, G. Laing, J. Wagner, S. Naisbitt and J. Salmond, "Validation of low-cost ozone measurement instruments suitable for use in an air-quality monitoring network," *Measurement Science and Technology*, vol. 24, no. 6, pp. 5803-5814, 2013.
- [29] U. Lerner, T. Yacobi, I. Levy, S. Moltchanov, T. Cole-Hunter and B. Fishbain, "The effect of ego-motion on environmental monitoring," *Science of the Total Environment*, vol. 533, pp. 8-16, 2015.
- [30] R. Piedrahita, Y. Xiang, N. Masson, J. Ortega, A. Collier, Y. Jiang, K. Li, R. Dick, Q. Lv, M. Hannigan and L. Shang, "The next generation of low-cost personal air quality sensors for quantitative exposure monitoring 7, 33," *Atmos. Meas. Tech.*, vol. 7, pp. 3325-3336, 2014.
- [31] N. Rabinovitch, M. Strand and E. Gelfand, "Particulate Levels Are Associated with Early Asthma Worsening in Children with Persistent Disease," *American Journal of Respiratory and Critical Care Medicine*, vol. 173, no. 10, pp. 1098-1105, 2006.
- [32] G. Kyrkilis, A. Chaloulakou and P. Kassomenos, "Development of an aggregate Air Quality Index for an urban Mediterranean agglomeration: Relation to potential health effects," *Environment International*, vol. 33, no. 5, pp. 670-676, 2007.
- [33] M. Kendall, Rank Correlation Methods, London: Charles Griffin & Company Limited, 1948.
- [34] W. Daniel, "Kendall's tau," in *Applied Nonparametric Statistics*, Boston, PWS-Kent, 1990, pp. 365-377.
- [35] J. Myers, A. Well and R. L. Jr, "Inference, assumptions, and power in multiple regression," in *Research Design and Statistical Analysis*, New-York, NY, USA, Lawrence Erlbaum, 2010, pp. 551-572.
- [36] B. Fishbain and E. Moreno-Centeno, "Self Calibrated Wireless Distributed Environmental Sensory Networks," *Scientific Reports*, in press.
- [37] D. Carslaw, S. Beevers, K. Ropkins and M. Bell, "Detecting and quantifying aircraft and other on-airport contributions to ambient nitrogen oxides in the vicinity of a large international airport," *Atmospheric Environment*, vol. 40, no. 28, pp. 5424-5434, 2006.

- [38] E. Westmoreland, N. Carslaw, D. Carslaw, A. Gillah and E. Bates, "Analysis of air quality within a street canyon using statistical and dispersion modelling techniques," *Atmospheric Environment*, vol. 41, no. 39, pp. 9195-9205, 2007.
- [39] A. Jones, R. Harrison and J. Baker, "The wind speed dependence of the concentrations of airborne particulate matter and Nox," *Atmospheric Environment*, vol. 44, no. 13, pp. 1682-1690, 2010.
- [40] D. Carslaw and K. Ropkins, "Openair d An R package for air quality data analysis," *Environmental Modelling & Software*, Vols. 27-28, pp. 52-61, 2012.
- [41] L. Yaroslavsky, *Theoretical Foundations of Digital Imaging Using MATLAB*, CRC Press, 2012.
- [42] P. Pokorná, J. Hovorka, M. Klán and P. K. Hopke, "Source apportionment of size resolved particulate matter at a European air pollution hot spot," *Science of the Total Environment*, vol. 502, no. 1, pp. 172-183, 2015.
- [43] R. Zhang, J. Jing, J. Tao, S.-C. Hsu, G. Wang, J. Cao, C. Lee, L. Zhu, Z. Chen and Y. a. o. Zhao, "Chemical characterization and source apportionment of PM 2.5 in Beijing: seasonal perspective," *Atmospheric Chemistry and Physics*, vol. 13, no. 14, pp. 7053-7074, 2013.
- [44] S. Arya, *Air pollution meteorology and dispersion*, New-York, NY, USA: Oxford University Press, 1999.
- [45] P. Zannetti, *Air pollution modeling: theories, computational methods and available software*, Springer Science & Business Media, 2013.
- [46] G. Kyrkilis, A. Chaloulakou and P. Kassomenos, "Development of an aggregate Air Quality Index for an urban Mediterranean agglomeration: Relation to potential health effects," *Environment International*, vol. 33, no. 5, pp. 670-676, 2007.
- [47] Y. Katznelson, "Ch. 1 - Fourier Series on T," in *An Introduction to Harmonic Analysis*, Cambridge, UK, Cambridge University Press, 2004, pp. 1-54.
- [48] M. Boas, *Mathematical Methods in the Physical Sciences*, Hoboken, NJ, USA: John Wiley & Sons , 2006.
- [49] W. Tsujita, A. Yoshino, H. Ishida and T. Moriizumi, "Gas sensor network for air-pollution monitoring," *Sensors Actuators B*, vol. 110, pp. 304-311, 2005.
- [50] AQMesh, UK, "AQMesh Website," July 2015. [Online]. Available: <http://www.aqmesh.com/>. [Accessed July 2015].

- [51] H. Yoshikado and M. Tsuchida, "High levels of winter air pollution under the influence of the urban heat island along the shore of Tokyo Bay," *Journal of Applied Meteorology*, vol. 35, no. 10, pp. 1804-1813, 1996.
- [52] J. H. Seinfeld and S. N. Pandis, *Atmospheric chemistry and physics: from air pollution to climate change*, John Wiley & Sons, 2016.
- [53] N. Smirnov, "Table for estimating the goodness of fit of empirical distributions," *Annals of Mathematical Statistics*, vol. 19, pp. 279-281, 1948.
- [54] T. B. Arnold and J. W. Emerson, "Nonparametric Goodness-of-Fit Tests for Discrete Null Distributions," *The R Journal*, vol. 3, no. 2, pp. 34-39, 2001.
- [55] N. Zumel and J. Mount, *Practical Data Science with R*, Shelter Island, NY, USA: Manning, 2014.
- [56] A. Tarantola, *Inverse problem theory and methods for model parameter estimation*, SIAM, 2005.

432

433