Feasibility of low-cost particle sensor types in long-term indoor air pollution health studies after repeated calibration, 2019–2021

Elle Anastasiou1, M. J. Ruzmyn Vilcassim3, John Adragna2, Emily Gill1, Albert Tovar1, Lorna E. Thorpe1 & Terry Gordon2

Previous studies have explored using calibrated low-cost particulate matter (PM) sensors, but important research gaps remain regarding long-term performance and reliability. Evaluate longitudinal performance of low-cost particle sensors by measuring sensor performance changes over 2 years of use. 51 low-cost particle sensors (Airbeam 1 N = 29; Airbeam 2 N = 22) were calibrated four times over a 2-year timeframe between 2019 and 2021. Cigarette smoke-specific calibration curves for Airbeam 1 and 2 PM sensors were created by directly comparing simultaneous 1-min readings of a Thermo Scientific Personal DataRAM PDR-1500 unit with a 2.5 µm inlet. Inter-sensor variability in calibration coefficient was high, particularly in Airbeam 1 sensors at study initiation. Calibration coefficients for both sensor types trended downwards over time to <1 at final calibration timepoint [Airbeam 1 Mean (SD) = 0.87 (0.20); Airbeam 2 Mean (SD) = 0.96 (0.27)]. We lost more Airbeam 1 sensors (N = 27 out of 56, failure rate 48.2%) than Airbeam 2 (N = 2 out of 24, failure rate 8.3%) due to electronics, battery, or data output issues. Evidence suggests degradation over time might depend more on particle sensor type, rather than individual usage. Repeated calibrations of low-cost particle sensors may increase confidence in reported PM levels in longitudinal indoor air pollution studies.

Abbreviations

PM  Particulate matter  
NAAQs  National ambient air quality standards  
SFH  Smoke-free housing  
EPA  Environmental Protection Agency  
PHA  Public Housing Authority

Studies of air pollution-associated health impacts often require measuring ambient concentrations of air pollutants. While monitoring of PM2.5 concentrations has contributed to understanding and reducing ambient PM2.5 to improve air quality standards, rigorous measurement of indoor air pollution remains a challenge.

Conventionally, the measurement of ambient PM1.5 concentrations requires either a labor-intensive gravimetric filter-based method with size-specific inlets, or sophisticated and manufacturer-calibrated real-time instruments. Such equipment is expensive and not readily portable, thus limiting the number of locations that can be sampled within a given time. Central monitoring networks that use advanced instruments utilizing gravimetry, light scattering, or beta attenuation have been mounted by states and federal agencies to address efforts to achieve federal national ambient air quality standards (NAAQS). Use of central monitoring is essential for monitoring

1Department of Population Health, New York University Grossman School of Medicine, 180 Madison Avenue, New York, NY 10016, USA. 2Department of Environmental Science, New York University Grossman School of Medicine, 341 East 25th Street, New York, NY 10010, USA. 3Department of Environmental Health Sciences, University of Alabama at Birmingham School of Public Health, Birmingham, AL 205-934-8927, USA. *email: terry.gordon@nyulangone.org
PM$_{2.5}$ exposures within the microenvironments of cities, in contrast, personal monitoring is considered the optimal approach to assess an individual’s exposure levels to PM$_{2.5}$.

Until recently, monitoring indoor settings at a high spatial and temporal resolution was impractical due to the cost, size, and expertise needed to operate monitoring equipment. Real-time personal monitoring or multi-location monitoring is, however, not a new concept. Innovations in air quality monitoring have addressed cost in the last decade. A combination of technological advancements (cheaper electronic boards and smaller light scattering sensors), public interest in air pollution, and the increased popularity of citizen science have resulted in the development and proliferation of low-cost PM$_{2.5}$ sensors and devices. These low-cost sensors have gained in popularity for a range of uses from home and personal monitoring to citizen science and to larger scale academic research.

The advantages of low-cost PM$_{2.5}$ sensors for research include: (1) deployment in large numbers to increase spatial and temporal coverage; (2) ease of use and maintenance; and (3) a battery power source that permits remote or portable use. In addition, they can be connected via Wi-Fi or Bluetooth technology to transmit data, sometimes in real time, to central servers and crowdsourcing platforms to share data and cover large geographic areas with extended spatial and temporal resolution. However, these simple, low-cost sensors have limitations and require routine testing and calibration prior to use in scientific studies. Much work has been done in recent years to address these limitations, and results have demonstrated that low-cost sensors generally have acceptable reliability but also technological limitations and inter-instrument variability. A key finding of many of these studies is that important research gaps remain regarding durability and the need for calibration of individual units prior to use for research.

Few studies, for example, have examined the performance of a network of low-cost sensors over an extended period. One such study showed that some PM$_{2.5}$ sensors were relatively stable over time when tested over a year, however that study focused on measurements in outdoor environments with concentrations ranging from 6 to 41 µg/m$^3$. Other short-term studies have demonstrated that careful calibration of low-cost sensors demonstrate their utility for indoor measurements of PM$_{2.5}$. In tandem, public housing authorities (PHAs) have been federally mandated to implement smoke-free housing (SFH) policies in their developments. Despite policy implementation in July 2018, there is still some evidence of cigarette smoking within New York City Housing Authority (NYCHA) developments. Stemming from a larger, quasi-experimental study evaluating the impact of SFH policies on secondhand smoke exposure in select NYCHA buildings, we utilized a network of low-cost sensors to evaluate indoor PM. This current analysis sought to assess whether rigorous calibration allows low-cost sensors to be used for indoor air quality measurements in the field for long periods of time without degradation and reliability. To achieve this objective, we repeatedly calibrated and utilized many low-cost first and second generation Airbeam PM$_{2.5}$ sensors, over a 2-year period, to assess PM$_{2.5}$ concentrations in urban high-rise buildings with a focus on measuring indoor tobacco smoke.

**Methods**

**Generation of calibration curves for cigarette smoke.** Cigarette smoke-specific calibration curves for the Airbeam 1 and 2 PM$_{2.5}$ sensors were created in a laboratory setting via the direct comparison of the output of the low-cost Airbeam sensors with simultaneous 1-min readings produced by a factory-calibrated Thermo Scientific Personal DataRAM PDR-1500 unit with a 2.5 µm inlet (Thermo Environmental Instruments, Waltham, MA). The PDR-1500 unit is a widely used instrument and shown to be reliable from previous studies. Over the course of the 2-year period, our low-cost sensors were calibrated four times using the same PDR-1500 unit.

To perform the calibration, 8-12 Airbeam units were placed into an airtight stainless-steel chamber, where temperature is room temperature and humidity matches the building’s at below 50%, with access ports permitting the introduction of cigarette smoke or HEPA filtered air. The PDR-1500 was connected to a sampling port for measuring the PM$_{2.5}$ concentrations inside the chamber. This instrument has both an inlet and outlet where tubes are connected to inject cigarette smoke into the chamber; the PDR-1500 was not placed inside the chamber to prevent contamination resulting from its enclosure with cigarette smoke. A smoking machine (Borgwaldt, Hamburg, Germany) was used to inject fresh mainstream cigarette smoke using 3R4F reference cigarettes into the chamber until the PDR-1500 registered a particle mass concentration greater than 1000 µg/m$^3$. A high concentration value such as 1000 µg/m$^3$ exceeds the upper limit for PM$_{2.5}$ values for both low-cost particle sensor types. Airbeam 1 and Airbeam 2 sensors have different saturation points at 80 µg/m$^3$ and 200 µg/m$^3$, respectively (i.e., the light scattering derived PM$_{2.5}$ output plateaus), ensuring the decreasing PM$_{2.5}$ calibration curve would begin above their detection ceiling (approximately 180 µg/m$^3$ and 800 µg/m$^3$, respectively). After cigarette smoke generation was stopped, the sample pump and internal filter of the PDR-1500 slowly removed cigarette smoke from the chamber which was replaced by HEPA-filtered room air. The resulting time-dependent decrease in PM$_{2.5}$ was used to develop the calibration curve. The start times of the Airbeam units and PDR-1500 particulate matter readings were synchronized, and the 1 min outputs were recorded beginning above the nominal upper detection limit and continued until the PDR-1500 values stabilized in the low single digit µg/m$^3$ range. Each run lasted approximately one hour.

Readings from each Airbeam (X-axis) were matched by synchronized timestamp with the corresponding values from the PDR-1500 (Y-axis). Using Excel, a unique calibration equation for each Airbeam unit was calculated by linear regression up to 80 µg/m$^3$ which was the expected upper limit for indoor PM$_{2.5}$. Polynomial regression models were also generated; however, the output of these models was linear up to 80 µg/m$^3$, which
Scientific Reports
www.nature.com/scientificreports/

The PM10 values from the Airbeam 2 sensors aligned closely with the corresponding values from the PDR-1500 sensors to approximate PM2.5 from the PDR-1500. Both sensor types use an algorithm based on an internal equation to generate PM output; Airbeam 1 sensors do not have the split for PM1, PM2.5 and PM10 values. The calibration coefficient for cigarette smoke was developed as a multiplication factor to correct the Airbeam PM2.5 output and calculated as:

\[
\text{Calibration Coefficient} = \text{slope of the PDR-1500 (Y-axis) vs Airbeam (X-axis) calibration curve}
\]

To assess the effect of particle composition on the calibration curve, the Airbeam devices were also calibrated using airborne particles in the NYC subway system. As in the cigarette smoke calibration procedure, the output of four Airbeam 1’s and four Airbeam 2’s was compared to the PDR 1500 PM2.5 output and a calibration coefficient was calculated for subway PM2.5.

**Field sampling periods.** We calibrated 51 low-cost particle sensors (Airbeam 1 generation N = 29; Airbeam 2 generation N = 22) at 4 different timepoints over a 2-year period spanning from 2019 to 2021. After each laboratory calibration, the Airbeam units were deployed in a large, natural experiment evaluating the impact of new smoke-free housing (SFH) policies on air quality in public housing units every 6 months18,30. Due to the onset of the COVID-19 pandemic, we were unable to perform Airbeam sensor calibration from April-September 2020. A technician-based calibration error for select Airbeam 2 sensors only, from December-March 2021, led to their exclusion from data analysis at that timepoint. The final calibration timepoint was collected for all 51 Airbeam sensors from May–September 2021 to obtain a final calibration coefficient.

**Data analysis.** We descriptively tabulated the mean (SD) calibration coefficients at four different 6-month timepoints over a 2-year period from 2019 to 2021 for the two different Airbeam sensor types. We performed independent t-tests to measure statistically significant differences in calibration coefficient means between particle sensor types, and characterized the between-and-within variability for calibration coefficient measurements. Because the light scattering properties of airborne particles are influenced by particle composition, we compared the mean (SD) calibration coefficients for cigarette smoke and subway PM2.5 using an independent t-test. Lastly, we used a difference-in-difference (DID) approach to compare within-group changes between Airbeam 1 and Airbeam 2 sensors across four different calibration timepoints. Regression models included fixed effects for particle sensor type (Airbeam 1 vs Airbeam 2 sensors) and data collection timepoints (12, 18, 30 and 36 months post-SFH policy implementation13). We adjusted for the clustering of individual Airbeam IDs and repeated measures overtime. Model-based mean differences with 95% confidence intervals were calculated for each particle sensor type over time. P-values were reported after implementation of the independent t-tests, with a significance level set at \( p < 0.05 \), using a two-sided test. All analyses were performed using SAS statistical software, version 9.4 (SAS institute).

We examined the individual time trends in calibration coefficient measurements for low-cost particle sensors over a 2-year period, grouped by particle sensor type (Supplemental Figure S1), and descriptively categorized all low-cost particle sensors that were taken out of circulation over the 2-year period (Supplemental Table S1). We then examined the correlation between the number of unique instances of use for individual Airbeam sensors, and their final calibration coefficients at the end of the 2-year period (Supplemental Table S2 and Supplemental Figure S2).

**Results**

**Sample characteristics.** We conducted a descriptive characterization of the mean (SD) calibration coefficients at four different timepoints over a 2-year timeframe from 2019 to 2021 (Table 1). At our first timepoint, our sample included a total of N = 56 Airbeam 1 sensors and N = 24 Airbeam 2 sensors. We observed more equipment failure over time in Airbeam 1 sensors (n = 27 out of 56, failure rate 48.2%) than in Airbeam 2 sensors (n = 2 out of 24, failure rate 8.3%). These equipment failures occurred for a variety of reasons including, but

| Time frame | Airbeam 1 (N = 29) | Airbeam 2 (N = 22) |
|------------|-------------------|-------------------|
|            | Mean   | SD     | Min   | Max   | Mean   | SD     | Min   | Max   | \( p \)-value |
| TimePoint 1| 1.43   | 0.44   | 0.96  | 2.64  | 1.56   | 0.26   | 1.23  | 2.07  | 0.21         |
| TimePoint 2| 1.14   | 0.22   | 0.65  | 1.69  | 1.58   | 0.26   | 0.97  | 2.02  | <0.0001      |
| TimePoint 3| 1.19   | 0.34   | 0.63  | 2.24  | –      | –      | –     | –     | –            |
| TimePoint 4| 0.87   | 0.20   | 0.55  | 1.23  | 0.96   | 0.27   | 0.74  | 1.89  | 0.2          |

Table 1. Descriptive characterization of calibration coefficient measurements among two low-cost particle sensor types over a two-year timeframe, 2019–2021. SD standard deviation.
not limited to cockroach infestations, not recording data properly (i.e., inconsistent relative humidity, temperature, or PM outputs), reading null values in PM measurements, and failure during calibration (Supplemental Table S1). As a result, our effective sample size decreased to $N = 37$ Airbeam 1 sensors and $N = 21$ Airbeam 2 sensors at the second timepoint, and $N = 29$ Airbeam 1 sensors and $N = 22$ Airbeam 2 sensors at the third and fourth timepoints. We thus restricted the analyses to the $N = 29$ Airbeam 1 sensors and $N = 22$ Airbeam 2 sensors available across all 4 calibration time points. We performed a secondary analysis to include all data points, both from units that performed well and from units that failed, and found no significant effect on outcome (see Supplement Table S3). The PM$_{2.5}$ concentration readout of Airbeam PM$_{2.5}$ sensors was less than that of the PDR-1500 reference instrument.

Between-and-within variability in calibration coefficients for low-cost particle sensor types. On an individual unit basis, we observed a high degree of inter-sensor variability in calibration coefficients across both low-cost particle sensor types over a 2-year timeframe (Fig. 1). There was a notable decline in Airbeam calibration coefficients consistent across both low-cost particle sensor types, with values trending downward to below one at the final calibration timepoint. Inter-monitor variability was high in Airbeam 1 sensors at the first calibration timepoint and in Airbeam 2 sensors at the fourth calibration timepoint. During the second calibration timepoint, the mean (standard deviation (SD)) calibration coefficient for Airbeam 1 sensors was lower compared to Airbeam 2 sensors [$M(\text{SD}) = 1.14(0.22)$ vs. $M(\text{SD}) = 1.58(0.26), p < 0.0001$] (Table 1). Because of the technical errors in the Airbeam 2 calibrations, a calibration coefficient mean was determined only for Airbeam 1 sensors ($1.19 (0.34)$) during the third calibration timepoint.

Least square mean differences in calibration coefficients for low-cost particle sensor types. We conducted a DID model approach for repeated measures and characterized the least square mean differences
Among Airbeam 1 sensors, the degree of inter-monitor change over time was statistically significant across all four time points, apart from change between the second and third timepoints \((-0.05 \ (−0.23, 0.14)\) \(p = 0.6\)). Among Airbeam 2 sensors, the degree of inter-monitor change over time was only statistically significant between the second and fourth timepoints \((0.62 \ (0.46, 0.78)\) \(p < 0.0001\) and between the first and fourth timepoints \((-0.60 \ (−0.76, −0.44)\), but not statistically significant between the first and second timepoints \((0.02 \ (−0.14, 0.18)\) \(p = 0.8\). We calculated the inter-monitor change over time among three timepoints only for Airbeam 2 sensors due to a calibration malfunction from December-March 2021.

Comparison of calibration coefficients for cigarette smoke versus subway particulate matter. Because particle composition can affect light scattering properties, we characterized the comparison of calibration coefficient mean differences by particulate matter source composition: an individual and pooled analysis across all units at a single timepoint. SD standard deviation.

Correlation between unique instances of use and final calibration coefficient for individual low-cost particle sensor types. To determine if sensor usage affected Airbeam output over time, we characterized the unique instances of use (i.e., the number of 7-day indoor sampling periods that a sensor was used), and the final calibration coefficient for all 51 individual Airbeam sensors (Supplemental Table S2). We examined the correlation between the number of 7-day indoor sampling periods that an individual sensor was used, and its final calibration coefficient at the fourth calibration timepoint (Supplemental Figure S2). We did not observe a strong correlation for Airbeam 1 sensors \(R^2 = 0.16\) for Airbeam 2 sensors \(R^2 = 0.09\). The slope of the curve for Airbeam 1 suggests that the more the sensors were used, the greater the deviation of its output from the PDR-1500’s output, while the Airbeam 2 curve suggested no change with an increase in usage.

Discussion

To our knowledge, this analysis is one of the first long-term longitudinal assessments of performance and reliability of low-cost particle sensors in measuring indoor tobacco smoking. We observed a high degree of inter-sensor variability across both particle sensor types, particularly in Airbeam 1 sensors at the study’s initiation. Change

| Effect | Mean difference (95% CI) | \(p\)-value | Mean difference (95% CI) | \(p\)-value |
|--------|--------------------------|-------------|--------------------------|-------------|
| Timepoint 1 to 2 | \(-0.29 \ (−0.42, −0.15)\) | < 0.0001 | \(0.02 \ (−0.14, 0.18)\) | 0.79 |
| Timepoint 1 to 3 | \(-0.24 \ (−0.41, −0.07)\) | 0.007 | - | - |
| Timepoint 1 to 4 | \(-0.56 \ (−0.70, −0.42)\) | < 0.0001 | \(-0.60 \ (−0.76, −0.44)\) | < 0.0001 |
| Timepoint 2 to 3 | \(-0.05 \ (−0.23, 0.14)\) | 0.63 | - | - |
| Timepoint 2 to 4 | \(0.28 \ (0.14, 0.42)\) | 0.0002 | \(0.62 \ (0.46, 0.78)\) | < 0.0001 |
| Timepoint 3 to 4 | \(0.32 \ (0.14, 0.50)\) | 0.0004 | - | - |

Table 2. Characterization of the least square mean differences for each low-cost particle sensor type over a two-year timeframe, 2019–2021.

| Individual unit | Calibration using cigarette smoke | Calibration using subway PM | Absolute mean difference |
|-----------------|----------------------------------|-----------------------------|-------------------------|
|                 | Mean (SD)                        | Mean (SD)                   |                         |
| 1               | 1.23 (0.84)                      | 0.39                        |                         |
| 2               | 1.30 (0.99)                      | 0.31                        |                         |
| 3               | 2.07 (0.82)                      | 1.25                        |                         |
| 4               | 0.97 (1.25)                      | 0.28                        |                         |
| 5               | 2.61 (1.33)                      | 1.28                        |                         |
| 6               | 2.63 (1.16)                      | 1.47                        |                         |
| 7               | 2.58 (2.04)                      | 0.54                        |                         |
| 8               | 0.93 (1.31)                      | 0.38                        |                         |
| Pooled analysis by calibration method | 1.79 (0.76) | 1.22 (0.39) | 0.08 |

Table 3. A comparison of calibration coefficient mean differences by particulate matter source composition: an individual and pooled analysis across all units at a single timepoint. SD standard deviation.
in calibration coefficients over time for individual Airbeam units was detected, suggesting a degradation of low-
cost particle sensors for longitudinal assessment (Supplemental Figure S1). There were also notable downward
trends in calibration coefficients over time, whereas the accompanying calibration coefficient for both Airbeam
1 and 2 sensors was below 1 at the final calibration timepoint. Findings lend support to the conclusion that the
routine calibration of individual Airbeam units might help to improve their utility and performance over time.
Overall, Airbeam 2 particle sensors fared better than Airbeam 1 sensors, suggesting greater durability of Airbeam
2 sensors for longitudinal assessment.

Our findings suggest that low-cost particle sensors might be differentially subjected to degradation, seen in
the greater loss of Airbeam 1 sensors than Airbeam 2 sensors over time. While two of these failures, and the loss
of units, resulted directly from the public housing environments (i.e., roach infestation), other failures were more
generally concerning for the use of non-calibrated low-cost particle sensors for longitudinal assessment of air
quality. Interestingly, we did not observe a strong correlation between the unique instances of field use of sen-
ors over the 2-year period and their final calibration coefficients measured at the fourth calibration timepoint,
suggesting that low-cost sensor degradation over time might be more contingent on particle sensor type, rather
than individual sensor usage.

Calibration coefficients differed modestly between cigarette smoke or subway PM (primarily combustion
products and iron-rich friction particles, respectively), suggesting that the light-scattering physics of these low-
cost particle sensors may slightly be affected by these two particle source types. Our finding, however, is limited
to two particular particle types, and further studies are needed to assess calibration across a range of particles
with different source-dependent compositions. Other researchers have observed that, in addition to particle
composition, the accuracy of PM$_{2.5}$ sensor output also depends upon particle size and humidity. Thus, low-cost
sensors require routine calibration in the laboratory with the PM$_{2.5}$ and environmental conditions of interest.

Our current analysis provides a robust assessment of the longitudinal utility of low-cost particle sensors.
Previous studies have measured the utility of low-cost particle sensors for PM monitoring where reference-
standard equipment is not available or feasible, and for improving the study of spatially localized airborne PM
concentrations. One study conducted in the United Kingdom evaluated the performance of four models of
low-cost PM sensors and examined inter-model performance across 19 different particle sensor units. Despite
similarities in the way each sensor type derived PM concentrations, the researchers found general agreement in
PM readings across sensor types. Another study evaluated the performance of two widely-used particle sen-
sors, the Plantower PMS A003 and Shinyei PPD42NS, and developed PM calibration models for seven different
metropolitan areas (i.e., Los Angeles, Chicago, New York, Baltimore, Minneapolis-St. Paul, Winston-Salem and
Seattle) using a sample of 72 sensors. The authors found that good calibration models were feasible only with the
Plantower PMS A003 model after running simulations for region-specific models. Another study found that a
Plantower PMS 1003 sensor provided reliable PM data outputs over a 13-month period. Our study extended
this time period to over 2 years of reliable output from Plantower PM sensors (albeit a different model), although
the reproducibility of the calibration coefficients varied by individual units over time. One of the largest programs
of low-cost sensor use is currently underway with the U.S. EPA’s AirNow network of low-cost PurpleAir sensors
for the nationwide monitoring of wildfire-generated PM. As demonstrated in our study of Airbeam sensors, the
PurpleAir sensors report PM levels that differ from more expensive and reliable monitoring instruments, but these offsets can be corrected by a ‘correction equation.’ The underlying design of the PurpleAir device is based on the fact that low-cost sensors may degrade over time and therefore the PurpleAir device evaluates individual sensor degradation by continually comparing
the output of two low-cost Plantower PM sensor units built into each monitoring device. As such, the EPA has
published guidelines on the use and performance testing of low-cost air pollution sensors. Without such corrections, caution is necessary regarding the reliability of low-cost PM sensors over time.

There were several limitations to our research. Overall, the PM output of each low-cost particle sensor differed
from the PM output of the widely used PDR-1500 which has an air flow regulation and infra-red laser that are far
more precise than what is available in the low-cost PM sensors, suggesting a potential for under- or over-
estimation of PM levels when calibration methods are not utilized. Over time, we experienced equipment failures
in a significant number of sensors, particularly the Airbeam 1 generation, thus reducing our effective sample size
in this calibration study. The results in our paper may fail to cover all the low-cost sensors and calibration of low-
cost PM sensors is imperative. Our routine calibration and inspection of low-cost particle sensors ensured careful
use for the long-term sampling of indoor tobacco smoke. Unfortunately, this type of calibration would likely be
challenging for many community groups or citizen science groups that may not have access to higher quality
PM monitors. There were also several strengths to our research. Our study provides a robust assessment of the
utility of low-cost particle sensors among a large number of a single brand of two generations of particle sensors
available for purchase and utilized in citizen science across the U.S. We compared the robustness of these two
low-cost Airbeam particle sensor types, as well as across two different calibration particle types. We restricted our
analysis to sensors that did not provide evidence of malfunction over time, and measured calibration coefficients
over a 2-year period, allowing for the assessment of the reliability of these particles for air quality monitoring.

Conclusions

We observed modest changes in calibration coefficient measurements over a 2-year timeframe among both
low-cost Airbeam particle sensor types, but in general the later generation Airbeam 2 model was more reliable,
suggesting that specific particle sensors may yield better longitudinal consistency. While we did observe a degree
of inter-monitor variability, changes in calibration coefficient measurements were relatively consistent across
Airbeam 1 and 2 sensors. Finally, while not significant, we observed a modest difference in calibration coefficients
when using cigarette smoke and subway PM as the calibration PM. As noted by our results and that of other researchers, low-cost PM sensors can provide reliable and consistent air quality data but regular calibration of the monitors is necessary to optimize their utility.

Data availability

Data generated or analyzed during this study can be found within the published article and its supplementary materials found in the appendices of this article.

Received: 19 October 2021; Accepted: 8 August 2022
Published online: 26 August 2022

References

1. Larkin, A. & Hystad, P. Towards personal exposures: how technology is changing air pollution and health research. *Carr Environ Health Rep* 4(4), 463–471. https://doi.org/10.1007/s40572-017-0163-y (2017).

2. Lim CC, Kim H, Vilcassim MIR, et al. Mapping urban air quality using mobile sampling with low-cost sensors and machine learning in Seoul, South Korea. *Environ Int.* 2019;131:105022. doi:https://doi.org/10.1016/j.envint.2019.105022

3. Morawski, L. et al. Applications of low-cost sensing technologies for air quality monitoring and exposure assessment: How far have they gone?. *Environ Int.* 2016;116, 286–299. https://doi.org/10.1016/j.envint.2018.04.018 (2018).

4. Reece S, Williams R, Colon M, et al. Spatial-Temporal Analysis of PM2.5 and NO2 Concentrations Collected Using Low-Cost Sensors in Penuelas, Puerto Rico. *Sensors (Basel).* 2018;18(12). doi:https://doi.org/10.3390/s18124314

5. Alfano B, Barretta L, Del Guadace A, et al. A review of low-cost particulate matter sensors from the developers’ perspectives. *Sensors (Basel).* 2020;20(23). doi:https://doi.org/10.3390/s20236819

6. Koehler, K. A. & Peters, T. M. New methods for personal exposure monitoring for airborne particles. *Sensors (Basel).* 2020;21(18)doi:https://doi.org/10.3390/s21181662

7. Zusman, M. *AirCasting Habitat Map,* 2021. https://www.habitatmap.org/airbeam.

8. Bulot FMJ, Johnston SJ, Basford PJ, et al. Long-term field comparison of multiple low-cost particulate matter sensors in an outdoor urban environment. *Sci Rep.* 2019;9(1):7497. doi:https://doi.org/10.1038/s41598-019-43716-3

9. Cho H, Baek Y. Practical particulate matter sensors and accurate calibration system using low-cost commercial sensors. *Sensors (Basel).* 2021;21(18)doi:https://doi.org/10.3390/s21181662

10. Holder AL, Mebust AK, Maghran LA, et al. Field evaluation of low-cost particulate matter sensors for measuring wildfire smoke. *Sensors (Basel).* 2020;20(17)doi:https://doi.org/10.3390/s20174796

11. Johnson, K. K., Bergin, M. H., Russell, A. G. & Hagler, G. S. W. Field test of several low-cost particulate matter sensors in high and low concentration urban environments. *Aerosol Air Qual Res.* 18(3), 565–578. https://doi.org/10.4209/aapqr.2017.10.0418 (2018).

12. Lee H, Kang J, Kim S, Im Y, Lee D. Long-term evaluation and calibration of low-cost particulate matter (PM) sensor. *Sensors (Basel).* 2020;20(13). doi:https://doi.org/10.3390/s20136317

13. Levy Zamora, M. et al. Field and laboratory evaluations of the low-cost plantower particulate matter sensor. *Environ Sci Technol.* 53(2), 838–849. doi:10.1021/acs.est.8b05174 (2019).

14. Mei H, Han P, Wang Y, et al. Field evaluation of low-cost particulate matter sensors in Beijing. *Sensors (Basel).* 2020;20(16). doi:https://doi.org/10.3390/s20164381

15. Liu, X. et al. Low-cost sensors as an alternative for long-term air quality monitoring, *Environ Res.* 185, 109438. https://doi.org/10.1016/j.envres.2020.109438 (2020).

16. Jiang Y, Zhu X, Chen C, et al. On-field test and data calibration of a low-cost sensor for fine particles exposure assessment. *Ecotoxicol Environ Saf.* 2021;211:111958. doi:https://doi.org/10.1016/j.ecosaf.2021.111958

17. Liu, M. et al. Using low-cost sensors to monitor indoor, outdoor, and personal ozone concentrations in Beijing, China. *Environ Sci Process Impacts.* 22(1), 131–143. https://doi.org/10.1039/c9em00377k (2020).

18. Thorpe LE, Anastasiou E, Wyzga K, et al. Evaluation of secondhand smoke exposure in New York City Public Housing after implementation of the 2018 Federal Smoke-Free Housing Policy. *JAMA Netw Open.* 2020;3(11):e2024385. doi:https://doi.org/10.1001/jamanetworkopen.2020.24385

19. Anastasiou E, Feinberg A, Tovar A, et al. Secondhand smoke exposure in public and private high-rise multiunit housing serving low-income residents in New York City prior to federal smoking ban in public housing, 2018. *Sci Total Environ.* 2020;704:135322. doi:https://doi.org/10.1016/j.scitotenv.2019.135322

20. U.S. Department of Housing and Urban Development (HUD). Taking matter into your own hands: About Airbeam. *AirCasting Habitat Map,* 2021. https://www.habitatmap.org/airbeam.

21. Wang, Z. et al. Comparison of real-time instruments and gravimetric method when measuring particulate matter in a residential building, *J Air Waste Manag Assoc.* 66(11), 1109–1120. https://doi.org/10.1080/10962247.2016.1201022 (2016).

22. Arif M, Parveen S. Carcinogenic effects of indoor black carbon and particulate matters (PM2.5 and PM10) in rural households of India. *Environ Sci Pollut Res Int.* 2021;28(21):2082–2096. doi:https://doi.org/10.1007/s11356-020-10668-5

23. Hallerman, A., Soussan, S. & Peters, T. M. Comparison of respirable mass concentrations measured by a personal dust monitor and a personal DataRAM to gravimetric measurements. *Ann Work Expo Health.* 62(1), 62–71. https://doi.org/10.1093/annweh/wxx083 (2017).

24. Marto, J. P., Zhang, J. & Schwab, J. J. Plume analysis from field evaluations of a portable air quality monitoring system. *J Air Waste Manag Assoc.* 71(1), 70–80. https://doi.org/10.1080/10962247.2020.1834010 (2021).

25. Oladejo, O. J. et al. An Assessment of Physicochemical characteristics of Awotan dumpsite in Ibadan, South western Nigeria. *Niger J Physiol Sci.* 35(1), 101–108 (2020).

26. Öljumí, O. O., Ana, G. R., Ogunseye, O. O. & Fabunmi, V. T. Air quality index from charcoal production sites, carboxyhemoglobin and lung function among occupationally exposed charcoal workers in South Western Nigeria. *Springerplus.* 5(1), 1546. https://doi.org/10.1186/s40064-016-3227-9 (2016).

27. Omí, T. M. & Ana, G. Ambient air pollution exposure and lung function assessment of filling station attendants in Ibadan, Nigeria. *Rev Environ Health.* 34(2), 211–218. https://doi.org/10.1515/reveh-2018-0027 (2019).

28. Singer, B. C. & Delp, W. W. Response of consumer and researcher grade indoor air quality monitors to residential sources of fine particles. *Indoor Air* 28(4), 624–639. https://doi.org/10.1111/iaa.12463 (2018).

29. Vilcassim MJ, Thurston GD, Polkier RE, Gordon T. Black carbon and particulate matter (PM2.5) concentrations in New York City's subway stations. *Environ Sci Technol.* 2014;48(24):14738–45. doi:https://doi.org/10.1021/es504295h

30. Cardozo RA, Feinberg A, Tovar A, et al. A protocol for measuring the impact of a smoke-free housing policy on indoor tobacco smoke exposure. *BMCPublic Health.* May 30 2019;19(1):666. doi:https://doi.org/10.1186/s12889-019-7043-3

31. PurpleAir, *PurpleAir: Real-time air quality monitoring everyone can use,* 2021; https://www2.purpleair.com/.

32. AirCasting Habitat Map, *Taking matter into your own hands: About Airbeam,* 2021. https://www.habitatmap.org/airbeam.
Acknowledgements
A.-M.F. and A.M. of New York City Housing Authority provided critical support for all aspects of data collection.

Author contributions
Drs. L.T. and T.G. had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis. Concept and design: L.T., T.G., E.A. Acquisition, analysis, or interpretation of data: L.T., E.A., A.T., E.G., T.G., J.A. Drafting of the manuscript: E.A., T.G., L.T., J.A., M.J.R.V. Critical revision of the manuscript for important intellectual content: All authors. Statistical analysis: T.G., E.A., R.V.

Funding
The work reported in this publication was supported by the National Cancer Institute of the National Institutes of Health under Award Number R01CA220591 and the National Heart Lung and Blood Institute of the National Institutes of Health under Award Number R01HL139239. The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Institutes of Health. The study protocol and procedures stemming from the R01CA220591 grant award were approved by the Institutional Review Board at the New York University School of Medicine on July 20, 2017; IRB number: S17-0968. In this current study stemming from our R01 grant, we did not involve human research subjects.

Competing interests
The authors declare no competing interests.

Additional information
Supplementary Information The online version contains supplementary material available at https://doi.org/10.1038/s41598-022-18200-0.

Correspondence and requests for materials should be addressed to T.G.

Reprints and permissions information is available at www.nature.com/reprints.

Publisher’s note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/.

© The Author(s) 2022