

# The particulate matter in the working environment of the digital printing machine detected by stationary and personal methods

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## Introduction



Particulate matter (PM) is the most significant pollutant produced by industrial activities. PM emitted during the digital printing process is potentially dangerous in the sense that it penetrates deep into the operator's lungs. To accurately assess the impact of these particles in the working environment, it's essential to determine their concentration and geometric characteristics. The analysis of PM is a complex process that requires the proposal of different analytical methods depending on the specific requirements and objectives of the research. The paper examines the similarities and differences between the concentrations of PM detected with spatial and personal sampling.

## Methods



### Digital printing environment

The research took place at the printing laboratory of the Department of Graphic Engineering and Design at the Faculty of Technical Sciences in Novi Sad, Serbia. PM mass concentration levels were measured near the ink jet digital printing machine, model LEC-540 (Roland, Japan).

### Analysis of particulate matter

#### Sensor sampler

PM particle mass concentration levels were continuously monitored using an optical particle counter OPC-N2 (Alphasense, United Kingdom) sensor sampler. The sensor device is a portable and precise PM monitor known for measuring PM mass concentrations ( $\mu\text{g}/\text{m}^3$ ) with high accuracy and real-time data collection.

#### Personal sampler

The EGO PLUS TT (Zambelli, Italy) personal sampler was used to collect PM particles with a diameter of less than  $10\ \mu\text{m}$ . The device consists of a display setting, a conical nozzle equipped with a filter, and a silicone hose. The membrane filters used were with a pore size of  $0.8\ \mu\text{m}$ , a diameter of  $25\ \text{mm}$  (effective diameter of  $22\ \text{mm}$ ), and an effective surface of  $380\ \text{mm}^2$ . The airflow rate was  $3.5\ \text{l}/\text{min}$ . The sampling time was 30 minutes, and measurements were taken within the printing technician's breathing zone, placed on the upper part of the chest.

The mass concentration levels (quantity) of PM particles with a diameter of less than  $10\ \mu\text{m}$  in the indoor air near the digital printing machine were determined with standard gravimetric measurement method according to the formula (1):

$$Q_{PM} = \frac{m_2 - m_1}{V} \cdot 10^6 \quad (1)$$

where:

- $Q_{PM}$  is the quantity of particles less than  $10\ \mu\text{m}$  ( $\mu\text{g}/\text{m}^3$ ),
- $m_1$  and  $m_2$  are the masses of the filter paper before and after sampling (g), respectively, and
- $V$  is the volume of air being passed through the device for 30 min ( $101.6\ \text{m}^3$ ).

## Results

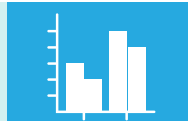


Figure illustrates the mass concentration levels of  $\text{PM}_{10}$ ,  $\text{PM}_{2.5}$ , and  $\text{PM}_1$  particles during 30 minutes of monitoring using a spatial sampling method with the sensor sampler, showing 8 values with a reading frequency of 4 minutes.

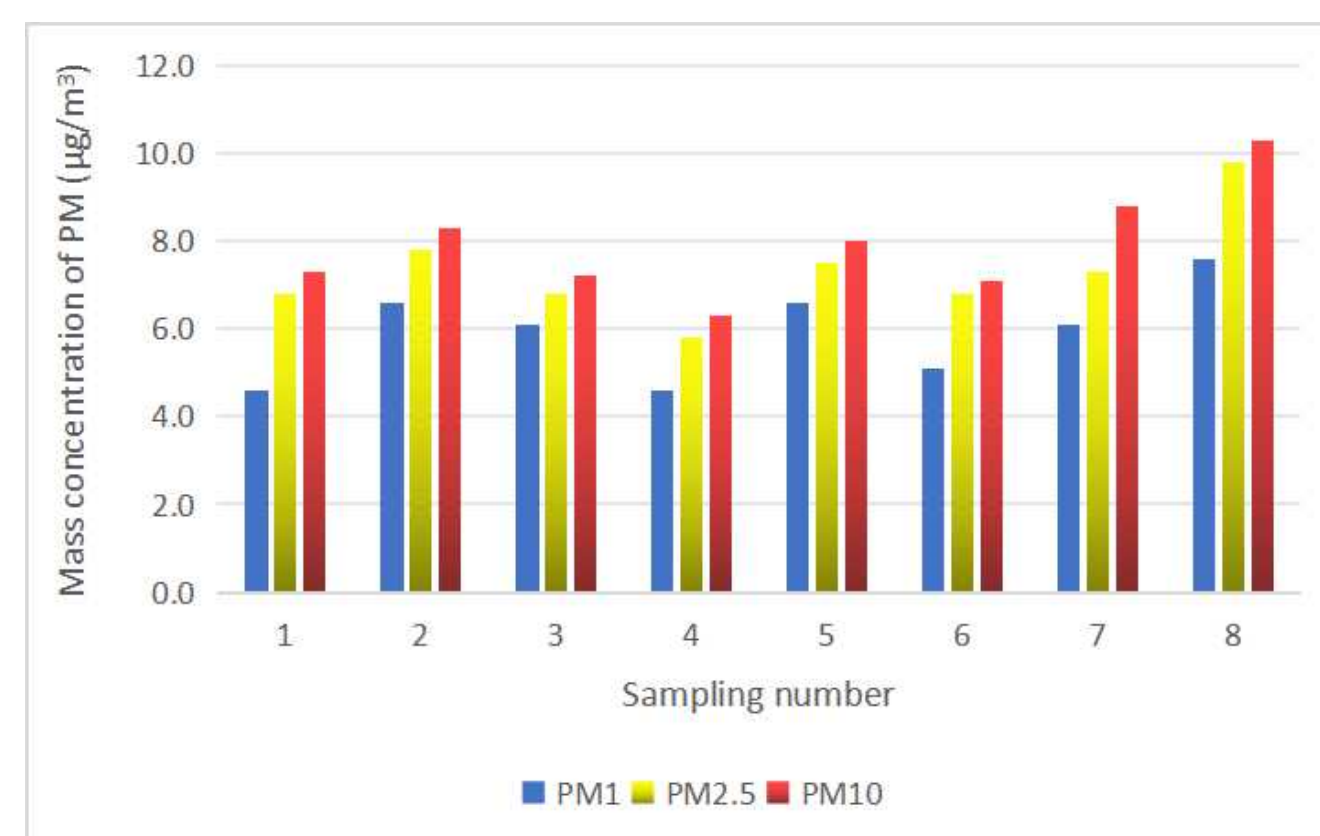


Figure 1

Concentration levels of  $\text{PM}_1$ ,  $\text{PM}_{2.5}$ , and  $\text{PM}_{10}$  particles detected by spatial sampling with sensor sampler

The spatial sampling method detected concentrations of  $\text{PM}_1$  particles in the range of  $4.6$  to  $7.6\ \mu\text{g}/\text{m}^3$  for 30 minutes. The concentrations did not grow exponentially, but an increase of 39.5% increase was observed from the 1<sup>st</sup> to the 8<sup>th</sup> sampling.

Over a 30-minute monitoring period using a sensor sampler,  $\text{PM}_{2.5}$  particle concentrations ranged from  $5.8$  to  $9.8\ \mu\text{g}/\text{m}^3$ . There was a 40.8% increase in concentrations from the lowest (detected in the 4<sup>th</sup> sampling) to the highest value (detected in the 8<sup>th</sup> sampling).

The mass concentrations of  $\text{PM}_{10}$  particles during the same 30-minute monitoring period ranged from  $6.3$  to  $10.3\ \mu\text{g}/\text{m}^3$ . An increase of 38.8% was observed from the lowest concentration (detected in the 4<sup>th</sup> sampling) to the highest (detected in the 8<sup>th</sup> sampling).

After using the personal sampling method, the mass of the filter paper before and after 30 minutes of sampling was  $0.0254$  and  $0.0374\ \text{g}$ , respectively. Based on the measured masses and the volume of air passing through the device in 30 min (formula 1), the quantity of PM particles with a diameter smaller than  $10\ \mu\text{m}$  was  $118.11\ \mu\text{g}/\text{m}^3$ .

The results indicate that the mass concentration obtained by personal sampling is 5.6 times higher (an 82% increase) than the cumulative value ( $21.1\ \mu\text{g}/\text{m}^3$ ) of  $\text{PM}_1$ ,  $\text{PM}_{2.5}$ , and  $\text{PM}_{10}$  detectable particles with a sensor sampler.

It's important to note that spatial and personal sampling methods may not always produce comparable results. The measurements using personal samplers tend to demonstrate a higher concentration than spatial ones. This has practical implications for air quality monitoring, as it suggests that personal samplers may be more effective in detecting pollutants in certain situations. Due to their light weight, size, and the fact that they do not require calibration, passive samplers are preferred over active air pump sensor samplers.

## Discussion / Conclusion



The results based on spatial sampling indicate that the detected mass concentrations decrease in the following order:  $\text{PM}_{10} > \text{PM}_{2.5} > \text{PM}_1$ . The concentrations of  $\text{PM}_{10}$  particles are higher than those of  $\text{PM}_1$  particles by 15.3% to 37.0% and more than 4.2% to 17% higher than  $\text{PM}_{2.5}$  particles. Additionally, concentrations of  $\text{PM}_{2.5}$  particles are higher than those of  $\text{PM}_1$  particles by 10.3% to 32.4%. Also, the cumulative value of  $\text{PM}_1$ ,  $\text{PM}_{2.5}$ , and  $\text{PM}_{10}$  detectable particles with a sensor sampler was 82% lower than the value detected by the gravimetric analysis after the personal sampler. In future research, the focus will be on improving methodology and incorporating new methods. For example, using scanning electron microscopy, particles can be characterized by their morphological and size distributions. Drying the filters to a constant mass will eliminate the influence of moisture, leading to more accurate results through gravimetric analysis. Analyzing and optimizing the printing process can enhance the result correlations. Identifying the best position for the devices is crucial for minimizing discrepancies between results obtained from different methods.

## ACKNOWLEDGMENTS

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