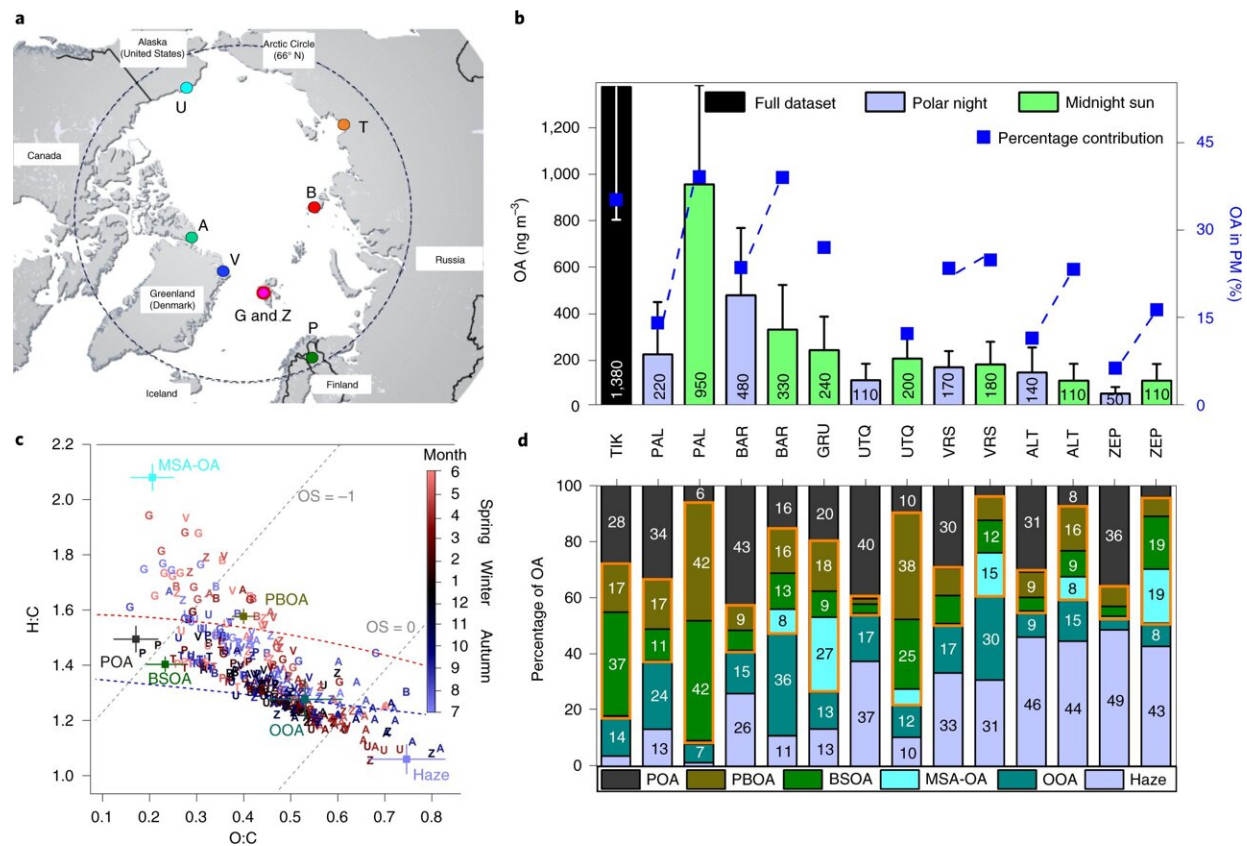


Scientists map Arctic aerosols to better understand regional warming

March 1 2022, by Clara Marc



Sites, OA factors and chemical characteristics. Credit: *Nature Geoscience* (2022). DOI: 10.1038/s41561-021-00891-1

Scientists at EPFL and the Paul Scherrer Institute (PSI) have studied the chemical composition and origin—whether natural or anthropogenic—of

aerosols in a region spanning from Russia to Canada. Their findings provide unique insights for helping researchers better understand climate change in the Arctic and design effective pollution-mitigation measures. The work was made possible thanks to the joint effort of scientists from three continents.

The [tiny particles](#) suspended in the air known as aerosols play an important role in heating and cooling our planet, but their effects still aren't fully understood. The particles can occur naturally, such as from volcanoes, forests and oceans, or be produced by human activity, such as fossil-fuel combustion and industrial manufacturing. They interact with solar radiation, either reflecting it back out into space and lowering the Earth's temperature, or absorbing it and raising the temperature. They are also essential for the formation of clouds, which similarly play a role in cooling off or warming up the planet by reflecting solar radiation out into space or re-emitting terrestrial radiation back down to the Earth. Cloud formation in the Arctic is particularly sensitive to aerosols.

To gain deeper insight into these mechanisms, scientists at ENAC's Extreme Environments Research Laboratory, headed by tenure-track assistant professor Dr. Julia Schmale, and the PSI's Laboratory of Atmospheric Chemistry, whose Research Laboratory Head is Dr. Imad El Haddad, analyzed samples taken from eight research stations across the Arctic over several years. The Arctic is a crucial region for understanding climate change because the temperature there is rising two to three times faster than the rest of the planet. "If we know what kind of aerosols exist in different areas and at different times of year, and what the origin and composition of those aerosols are, we will have a better grasp of how they contribute to climate change," says Schmale. "That will also help us design more targeted measures to reduce pollution." The study was led by Vaios Moschos as part of his Ph.D. thesis, supervised jointly by Schmale and El Haddad.

Anthropogenic in the winter and natural in the summer

In a first study, Moschos et al. looked specifically at organic aerosols. Scientists still have little data on these aerosols even though they make up nearly 50% of total particulate matter. The researchers in this study analyzed the [chemical composition](#) of samples taken in the Arctic and found that, in the winter, most of those aerosols come from human activity. They attribute this to the Arctic haze that occurs each year when emissions from oil extraction and mining operations in North America, Eastern Europe and Russia are carried to the Arctic and trapped there during the winter.

On the other hand, the study found that most organic aerosols in the summer come from natural sources. That's because the transport of anthropogenic aerosols from mid-latitudes to the Arctic is diminished during the warmer months, and the high latitude emission rate of biogenic aerosols or their precursors rises. "We didn't expect to see so much naturally occurring organic aerosols," says Schmale. "These particles come from boreal forests as well as phytoplankton, a micro-organism that lives in oceans. Here we might see a consequence of global warming in the future—as forests expand northwards and the permafrost thaws more organic molecules can be released from land, and as sea ice retreats, more open ocean leaves space for microbial emissions."

Mitigation is now possible

In another study, the EPFL and PSI scientists used the same samples but analyzed the composition and origin of all the aerosols, both organic and inorganic. They found that the inorganic aerosols included [black carbon](#), sulfate and sea salt; black carbon is of particular concern to the scientific

community because it absorbs [solar radiation](#) and contributes to global warming. "We knew that black carbon emissions were high in regions with oil and gas extraction facilities, but we didn't have collocated pan-Arctic measurements to understand how large their circle of influence is," says Schmale. "Thanks to the data collected in this study, we were able to map black carbon concentrations and origins in each Arctic region throughout the year and recommend the most appropriate measures to take."

The scientists were able to perform the studies thanks to a unique joint effort bringing together scientists from Canada, Denmark, Finland, France, Germany, Greece, India, Italy, Norway, Russia, Slovenia and the US. The eight research stations at which samples were collected (see list below) are run by research groups from various countries. The samples were analyzed at the two labs in Switzerland. El Haddad explains: "Analyzing organic aerosols requires mass spectrometers, which are expensive, sophisticated instruments, along with trained experts. That's one reason why we still have little data from the Arctic on this subject. Our lab is at the forefront of research on organic aerosols and their origin."

Samples were collected at the following [research stations](#):

- Alert, Canada
- Baranova, Russia
- Gruvebadet, Norway
- Pallas, Finland
- Tiksi, Russia
- Utqiagvik, U.S.
- Villum, Greenland
- Zeppelin, Norway

The research was published in *Environmental Research Letters* and

Nature Geoscience.

More information: Vaios Moschos et al, Elucidating the present-day chemical composition, seasonality and source regions of climate-relevant aerosols across the Arctic land surface, *Environmental Research Letters* (2022). [DOI: 10.1088/1748-9326/ac444b](https://doi.org/10.1088/1748-9326/ac444b)

Vaios Moschos et al, Equal abundance of summertime natural and wintertime anthropogenic Arctic organic aerosols, *Nature Geoscience* (2022). [DOI: 10.1038/s41561-021-00891-1](https://doi.org/10.1038/s41561-021-00891-1)

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