

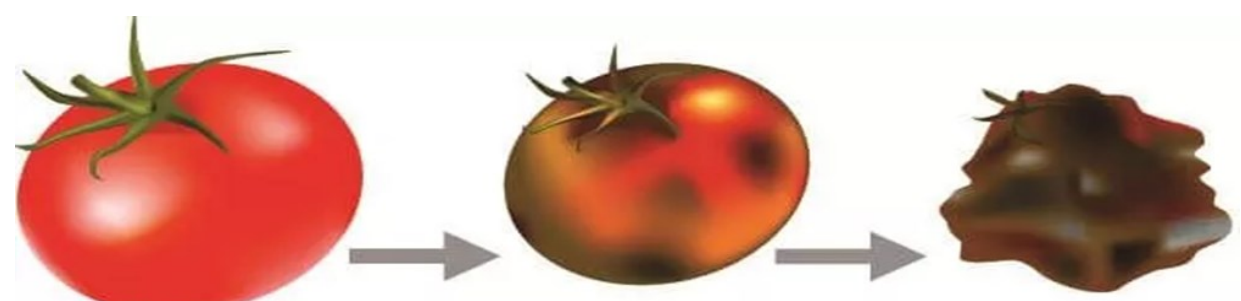
Long-term measurements of aerosol **Oxidative Potential** in an urban site, Athens, Greece

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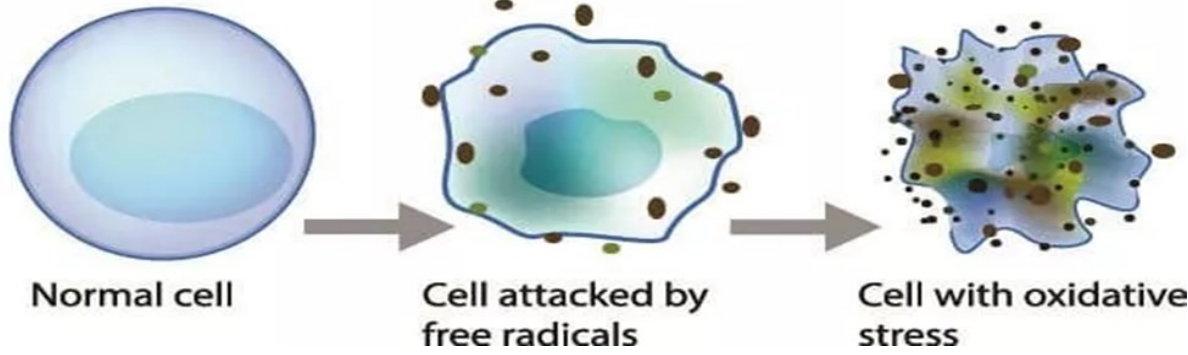
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<https://www.noa.gr>



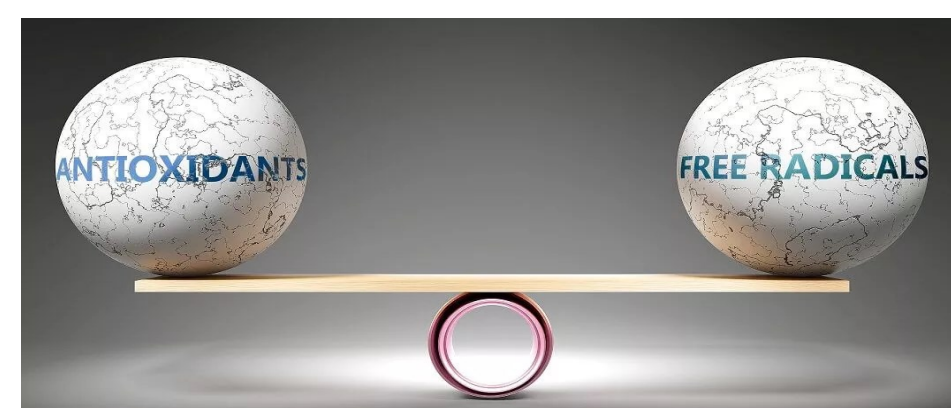
OXIDATIVE STRESS



<https://biologydictionary.net/oxidative-stress/>

Oxygen carrying molecules with an uneven number of electrons → **Free radicals**

Antioxidants → cause the free radical to stabilize and become less reactive.



When there is an imbalance, it leads to **Oxidative Stress** → **Cell damage**

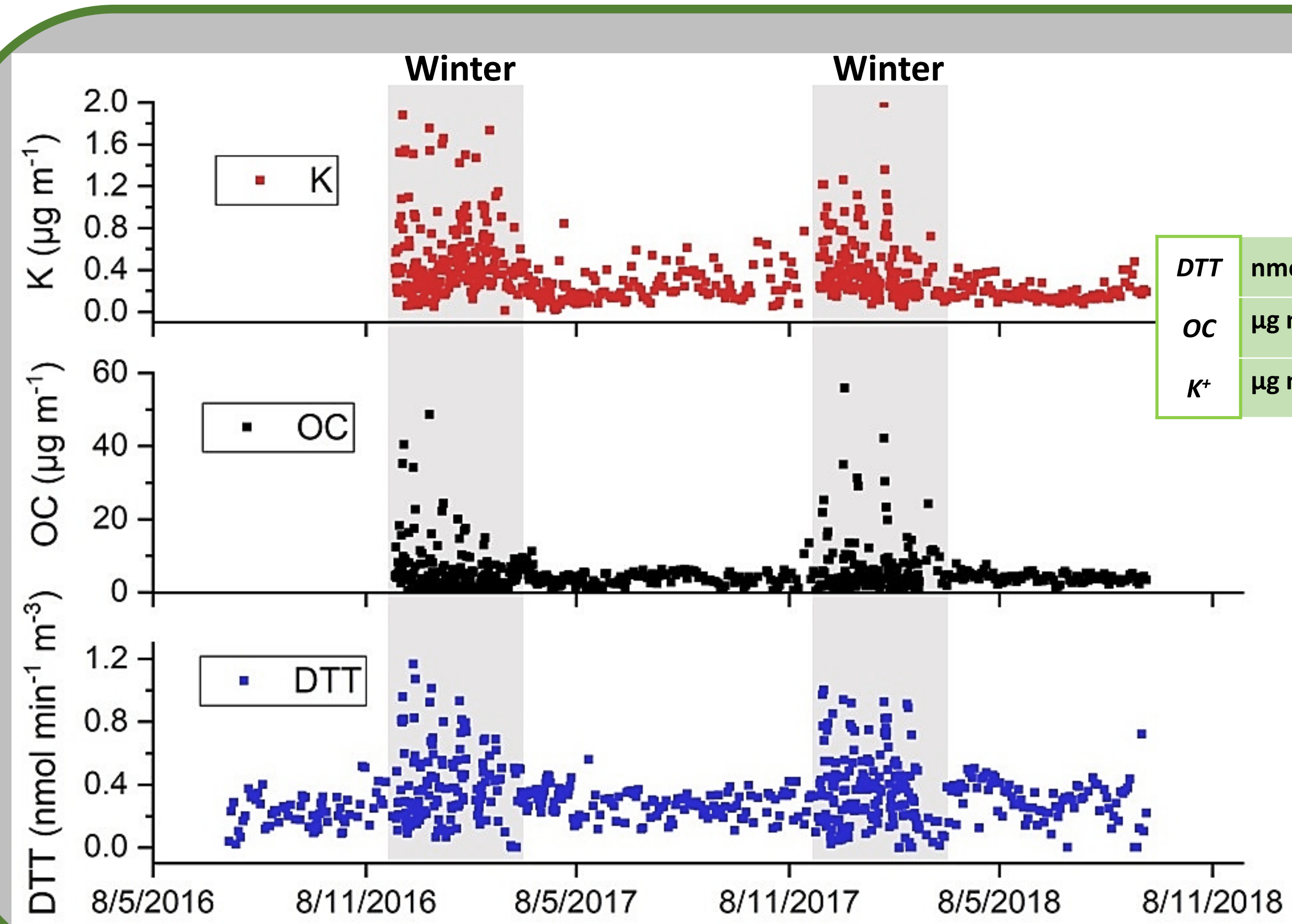
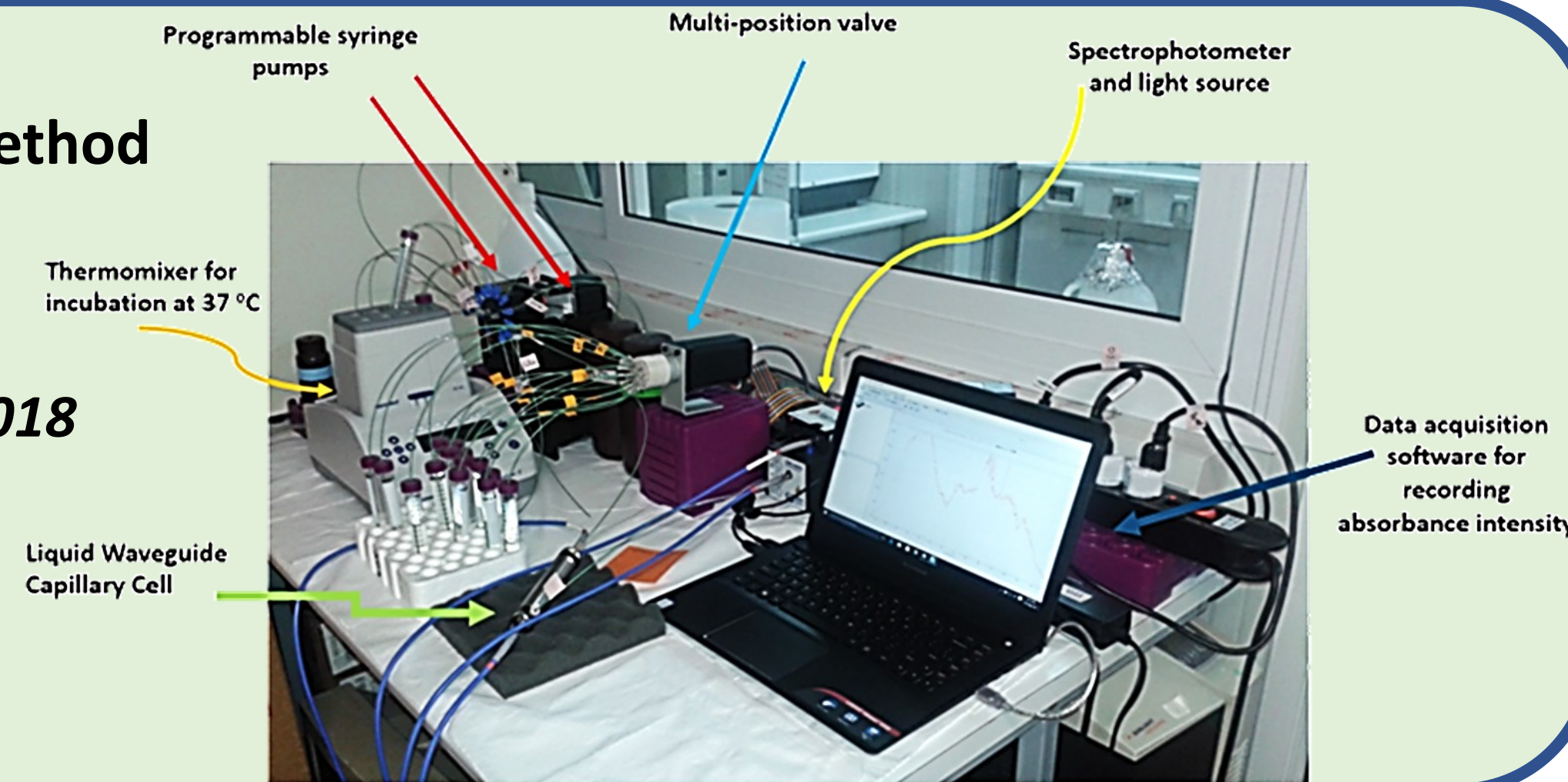
Oxidative Potential (OP)

Semi automated system → DTT assay method

Fang et al., 2015, Atmos. Meas. Tech., 8: 471-482.
Paraskevopoulou et al., 2019, Atm. Env., 206: 183-196

City center Sampling: July 2016 – July 2018

- PM_{2.5} mass concentrations
- Anions and Cations
- Organic and Elemental Carbon
- Trace metals and Elements

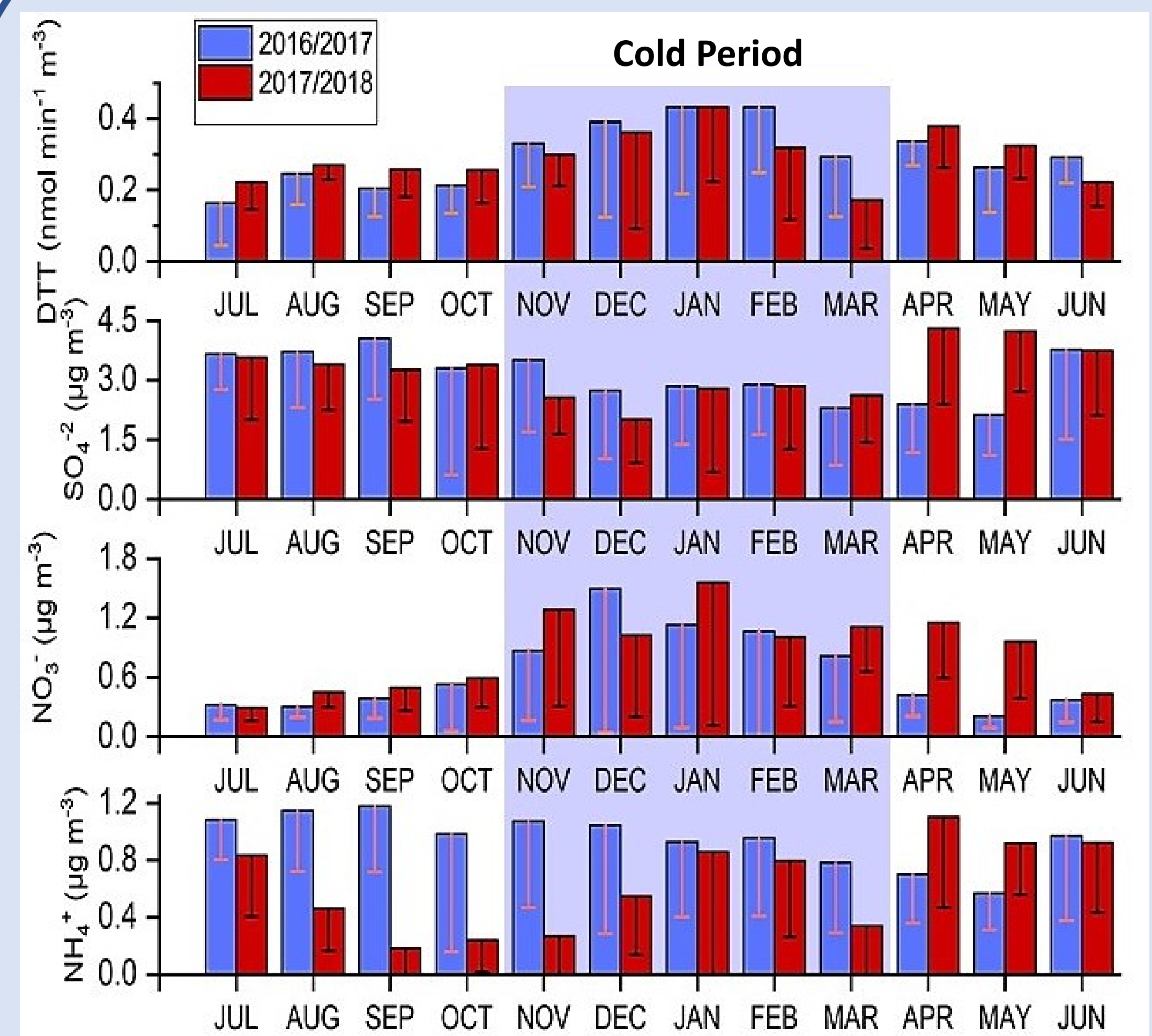


Two years daily variability

	Average	Range
DTT	0.33	1.16 - 0.01
OC	5.11	55.8 - 0.20
K ⁺	0.38	1.99 - 0.01

During winter, DDTv presents covariance with K⁺ and OC, suggesting that these components could possess considerable OP; while a link to biomass burning sources is highlighted.

Monthly variations



During the **coldest period** of the year biomass burning dominates, while in the **warmest months**, there is significantly higher contribution of secondary aerosol processing.

In the **warmest period** - although we cannot exclude the organic part of the aerosol - it appears that long-range transport of inorganic aerosol dominates in the OP of fine particulate matter.

	DTTv	K	OC	EC	NH ₄ ⁺	NO ₃ ⁻	SO ₄ ²⁻	Al	Cr	Co	Fe	Cu	Ni
Cold period	R ²	0.35	0.41	0.42	0.59	0.48	0.24	0.04	0.30	0.23	0.30	0.27	0.10
	n	284	290	290	282	284	284	95	95	58	90	95	93
Warm period	R ²	0.07	0.34	0.36	0.48	0.03	0.48	0.21	0.07	0.12	0.38	0.35	0.10
	n	82	527	527	72	72	72	35	35	37	35	35	35

Regression Analysis

Oxidative Potential vs. Chemical composition of aerosol

Better Pearson correlation in the **winter-time** → K⁺, NO₃⁻, Cr and Co
In **summer** → SO₄²⁻ and Al → better correlations compared to the coldest period of the year.
Long-range transport of inorganic aerosol → dominates in the OP of fine particulate matter during summer, along with the presence of Fe and Cu; while organic aerosol as well appears to contribute to a lesser extent.

ACKNOWLEDGMENTS

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CONCLUSIONS

- ❖ OP higher values during wintertime → wood burning dominant source of aerosol → combustion major source of water-soluble OP, both as primary and secondary emission
- ❖ Winter regression analysis → toxicity of aerosol is significantly associated with OC and NO₃⁻ → Once again, significant impact of combustion, both as primary and secondary product emission.
- ❖ During summer → an impact of long-range transportation and water-soluble metals in aerosol toxicity is shown.
- ❖ Combination of various PM chemical parameters → scarce identification of various aerosol OP sources → temporal basis, in the area of Eastern Mediterranean.