Triple Isotopes (δ¹³C, δ²H, and Δ¹⁴C) Compositions and Source Apportionment of Atmospheric Naphthalene: A Key Surrogate of Intermediate-Volatility Organic Compounds (IVOCs)

Tiangang Tang^{1,2}, Zhineng Cheng^{1*}, Gan Zhang^{1*}

¹State Key Laboratory of Organic Geochemistry, Guangzhou Institute of Geochemistry, Chinese Academy of Sciences, Guangzhou, 510640, P. R. China (*correspondence: zhanggan@gig.ac.cn; chengzhineng@gig.ac.cn)

²University of Chinese Academy of Sciences, Beijing 100049, P. R. China.

Widely occurring in the atmosphere, naphthalene (NAP) is worth concern not only owing to its carcinogenicity, but also as an important precursor of atmospheric SOA [1]. However, its source attributions are not explicit. We conducted source apportionment of atmospheric NAP using a triple-isotope (δ^{13} C, δ^{2} H, and Δ^{14} C) technique combined with a Bayesian model in the Beijing-Tianjin-Hebei (BTH) region, China [2].

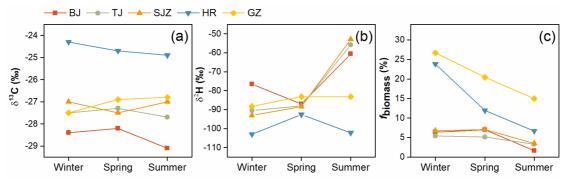


Figure 1. Temporal-spatial distribution of stable carbon (a), hydrogen (b) isotope compositions and fraction modern (c) of atmospheric vapor-phase naphthalene. Beijing (BJ), Tianjin (TJ), Shijiazhuang (SJZ), Huairou (HR) and Guangzhou (GZ).

At the urban sites (BJ, TJ, and SJZ), stable carbon ($-27.7 \pm 0.7 \%$) and radiocarbon ($-944.0 \pm 20.4\%$) isotope compositions of NAP did not exhibit significant seasonal variation, but the deuterium system showed a relatively more ²H-depleted signature in winter ($-86.7 \pm 8.9 \%$) compared to that in summer ($-56.4 \% \pm 3.9 \%$). Radiocarbon signatures indicated that 95.1 ± 1.8 % of NAP was emitted from fossil sources in these cities. The Bayesian model results indicated that the source contribution of the BTH urban sites had a similar pattern. The contribution of liquid fossil combustion was highest (46.7 ± 2.6%), followed by coal high-temperature combustion ($26.8 \pm 7.1\%$), coal low-temperature combustion ($18.9 \pm 6.4\%$) and biomass burning ($7.6 \pm 3.1\%$). At the suburban site, the contribution of coal low-temperature combustion could reach 70.1 ± 6.4 %. The triple-isotopes based approach provides a top-down constraint on the sources of atmospheric NAP, and could be further applied to other IVOCs in the ambient atmosphere.

- [1] Huang et al. (2019) Environ. Sci. Technol. 53, (3), 1235-1244.
- [2] Bosch et al. (2015) Environ. Sci. Technol. 49, (13), 7657-7665.