

How does smoke change as it is carried away from the source?

Emily Fischer

(evf@atmos.colostate.edu)

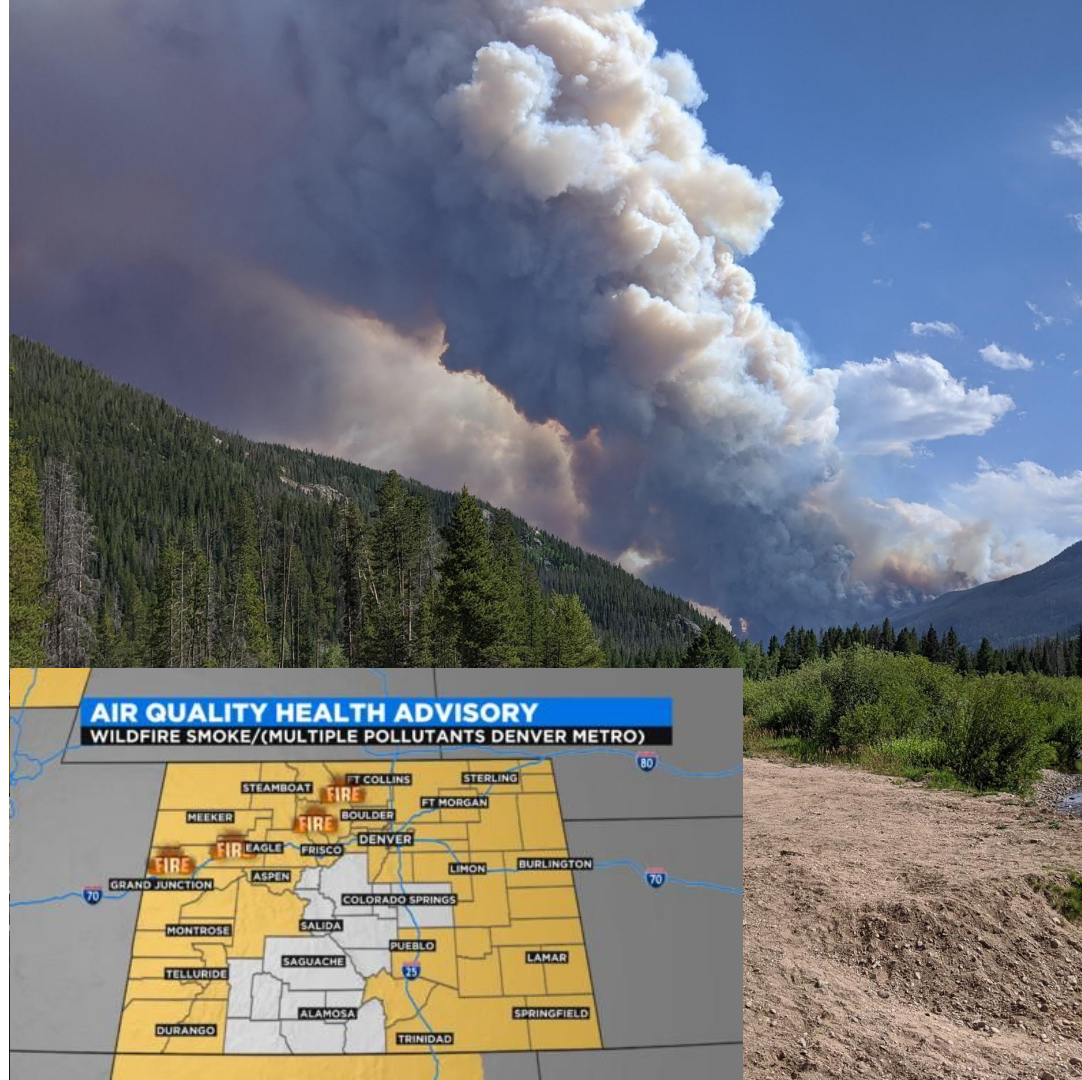
Thanks to these colleagues for their contributions to these remarks:

Jeffrey Pierce, Zachary Decker,
Steve Brown, and Brett Palm

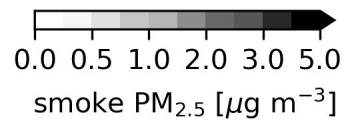
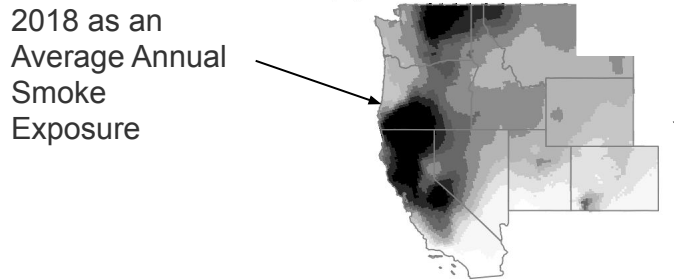
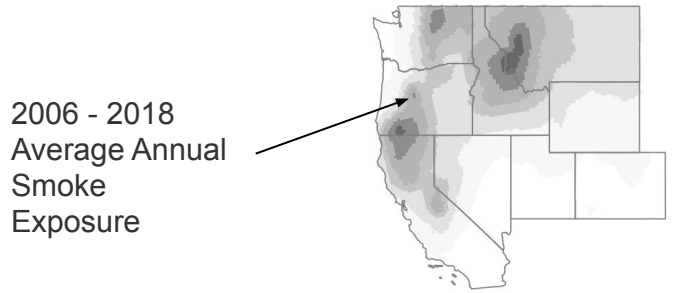
Thanks to the larger WE-CAN team.



NSF Award Numbers: AGS-1650786 (CSU), AGS-1650275 (U Montana), AGS-1650493 (U Wyoming), AGS-1650288 (CU-Boulder), AGS-1652688 (U Washington)
NOAA Award Number: NA17OAR4310010

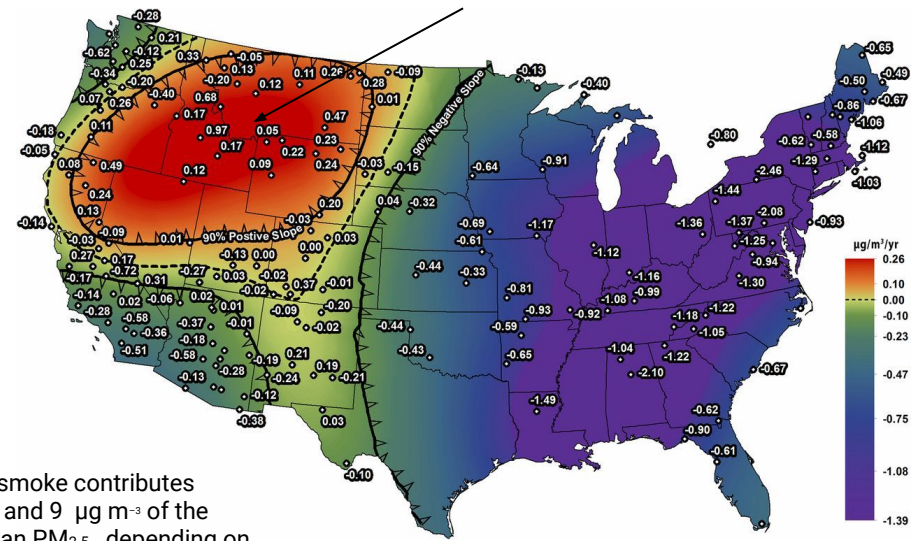


When we think of smoke, we first think of $PM_{2.5}$. Smoke makes the largest contribution to $PM_{2.5}$ in the Pacific Northwest and directly downwind.



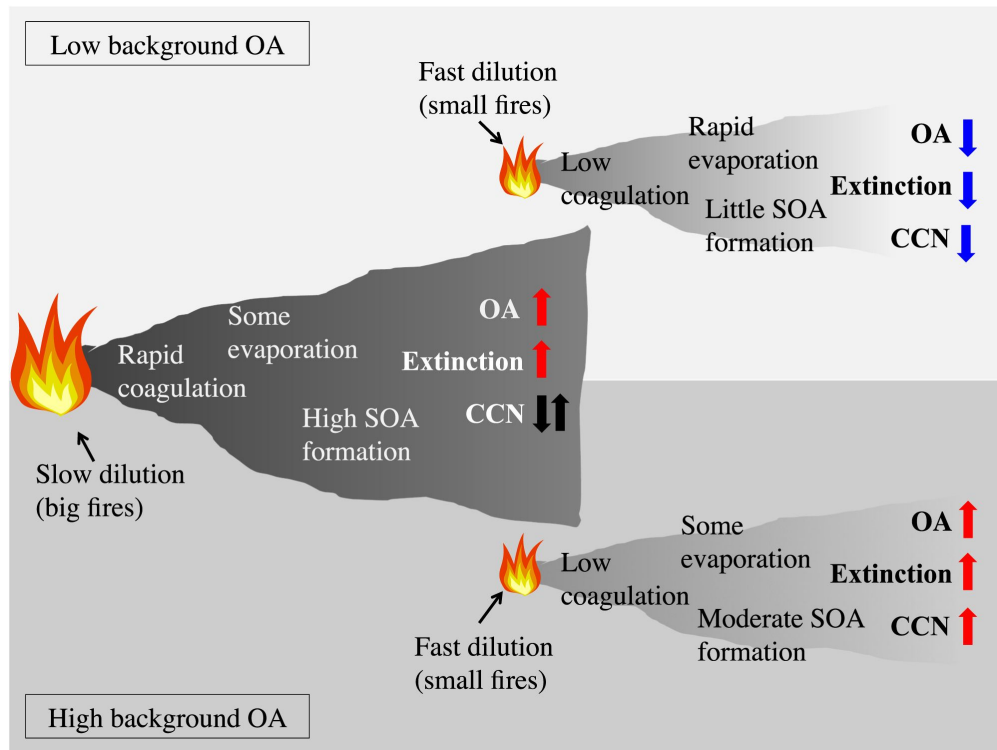
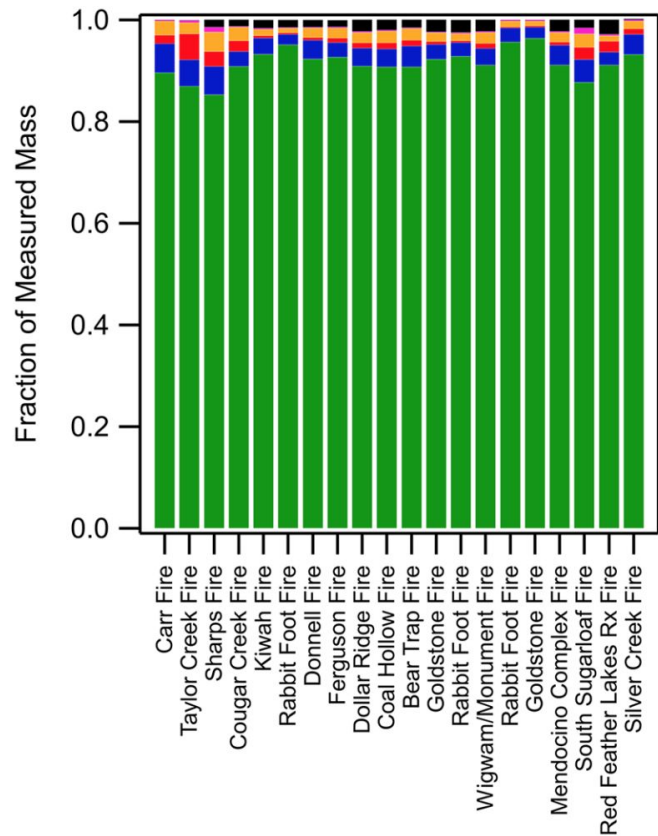
In the PNW smoke contributes between ~ 2 and $9\ \mu g\ m^{-3}$ of the summer-mean $PM_{2.5}$, depending on the year. In a high-fire year, this is $\sim 70\%$ of the $PM_{2.5}$.

The positive trend in 98th quantile $PM_{2.5}$ is due to wildfire activity

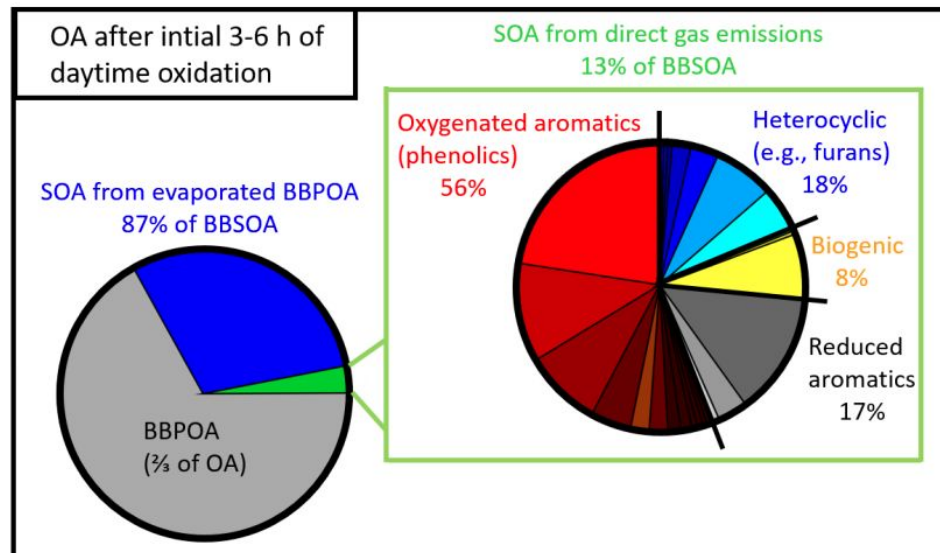
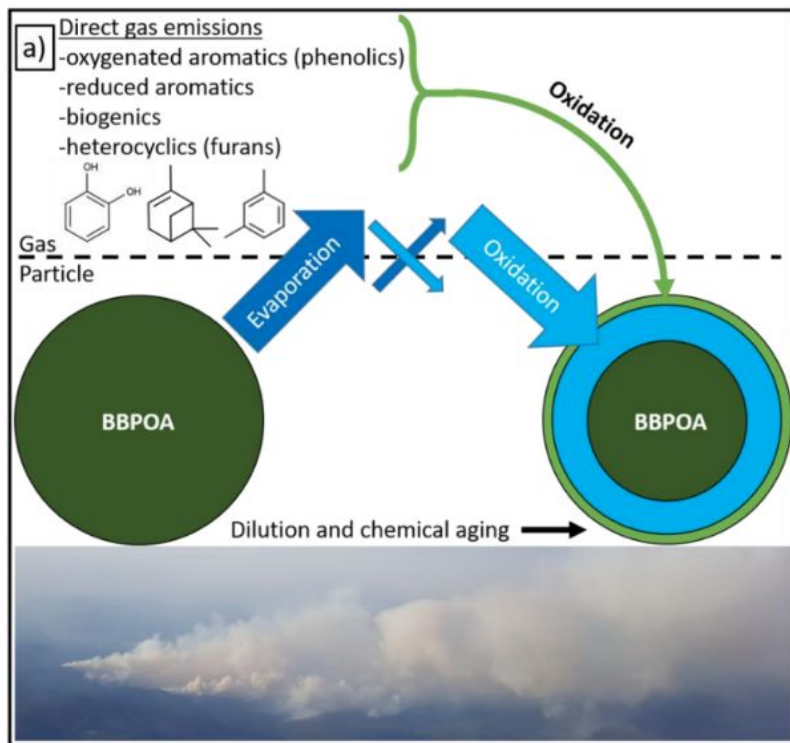


O'Dell et al. (2020); McClure et al. (2018)

Organic aerosols (OA) dominate fine PM mass. Recent work shows the interplay between emissions and subsequent chemical and physical transformations for OA.

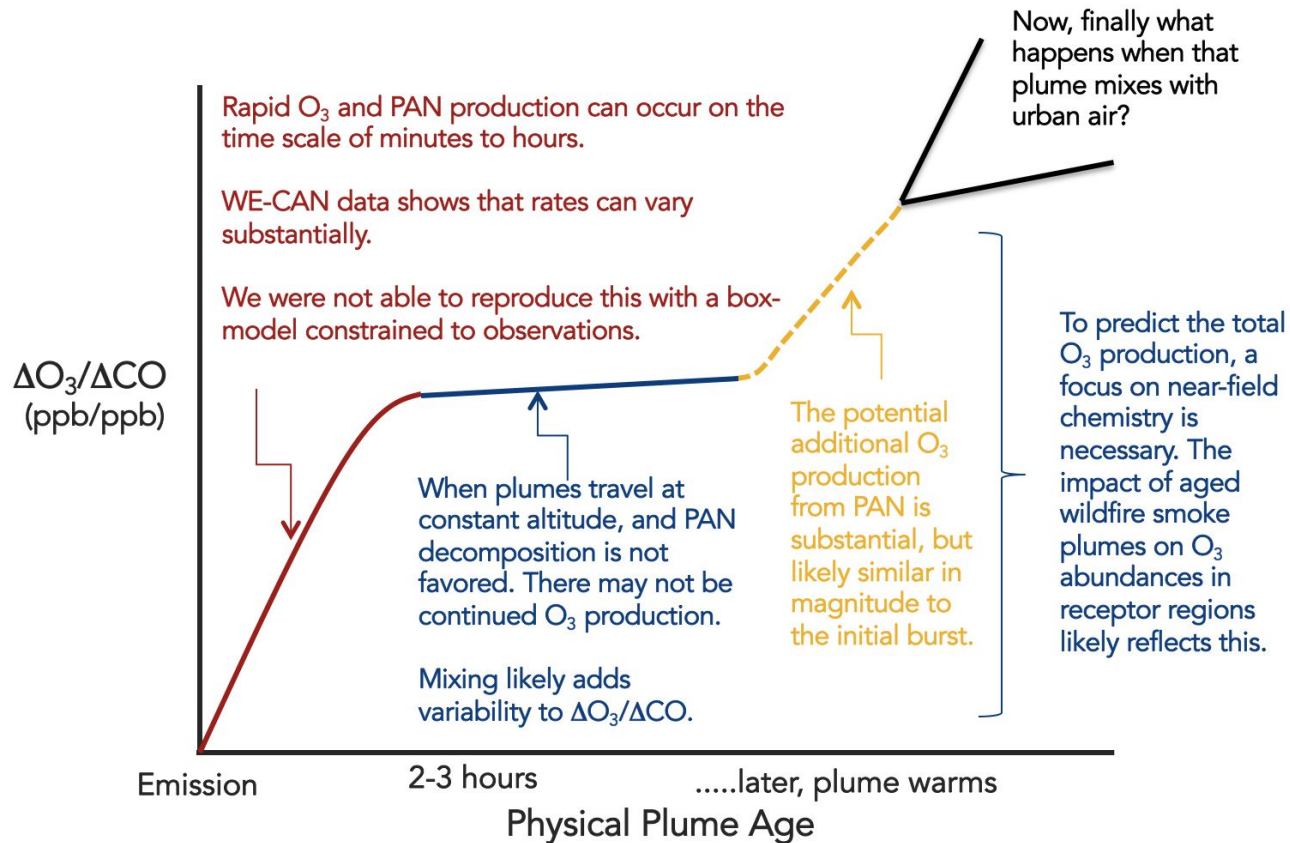


Phenolics are important gas-phase emissions (small influence on OA mass, important for absorption); evaporated POA is the dominant source for SOA.

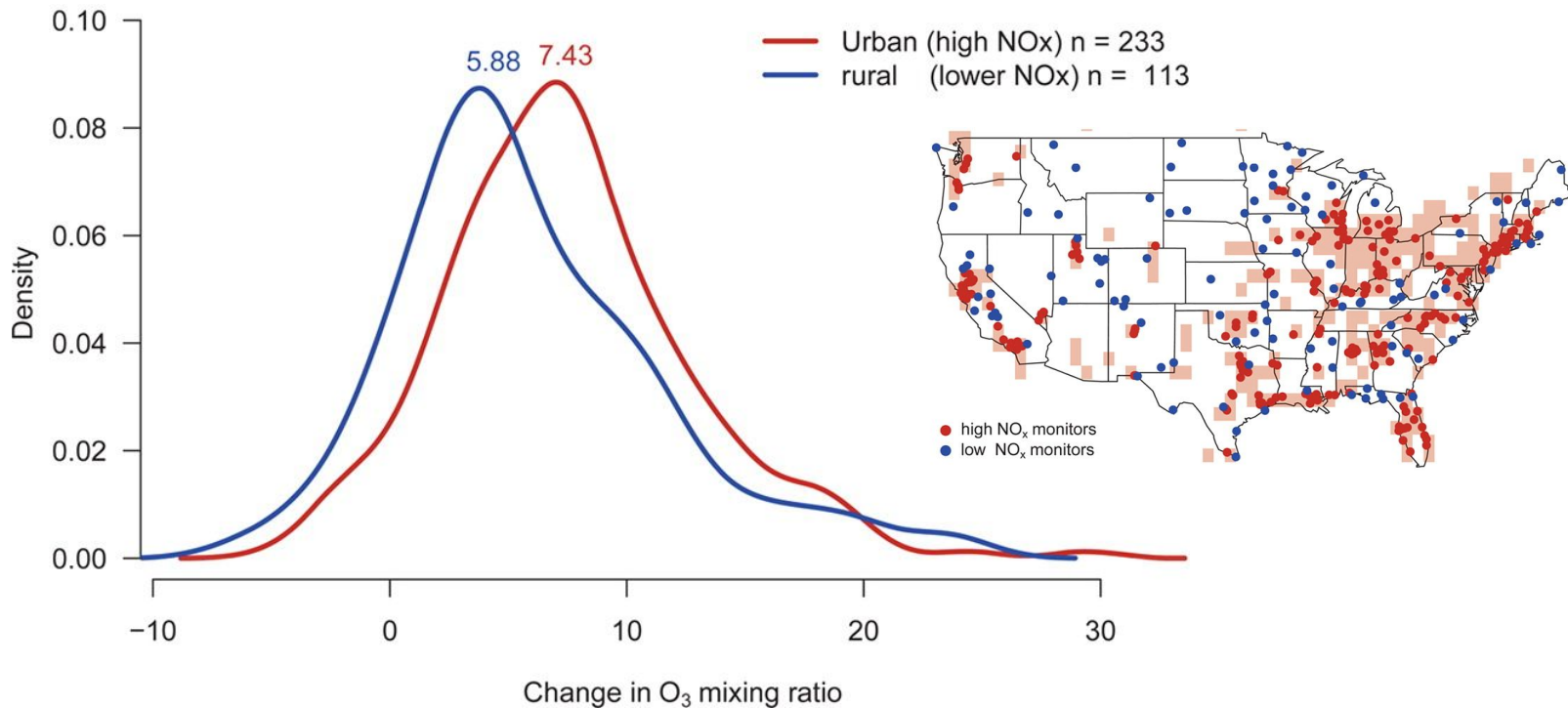


This study covers the period when large wildfire plumes dilute by a factor of 5 - 10 during the day.

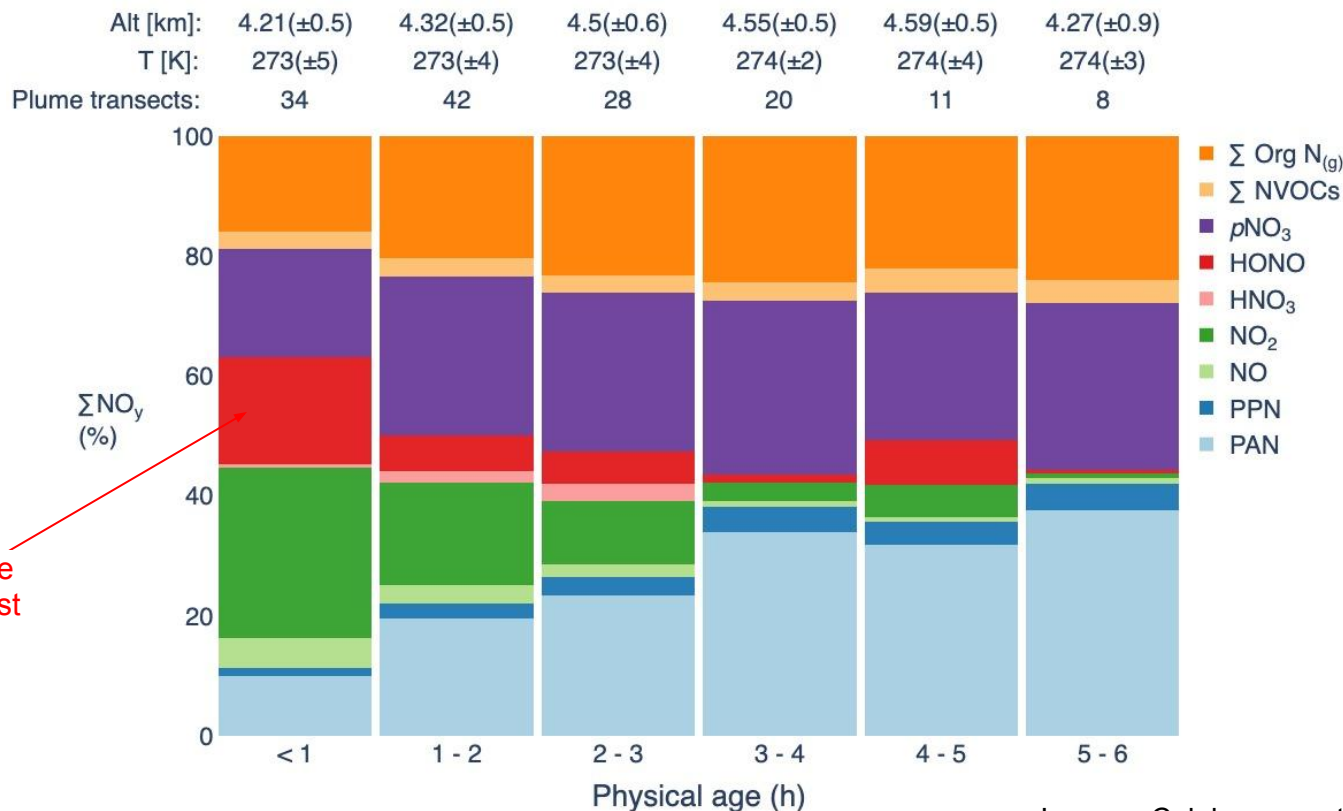
O₃ is often produced in smoke plumes, but there are different timescales and environmental conditions to consider.



Wildfire smoke impacts O_3 in both urban and rural locations. Across the U.S. the average increase in O_3 on a smoke-impacted day is ~ 6 ppbv.

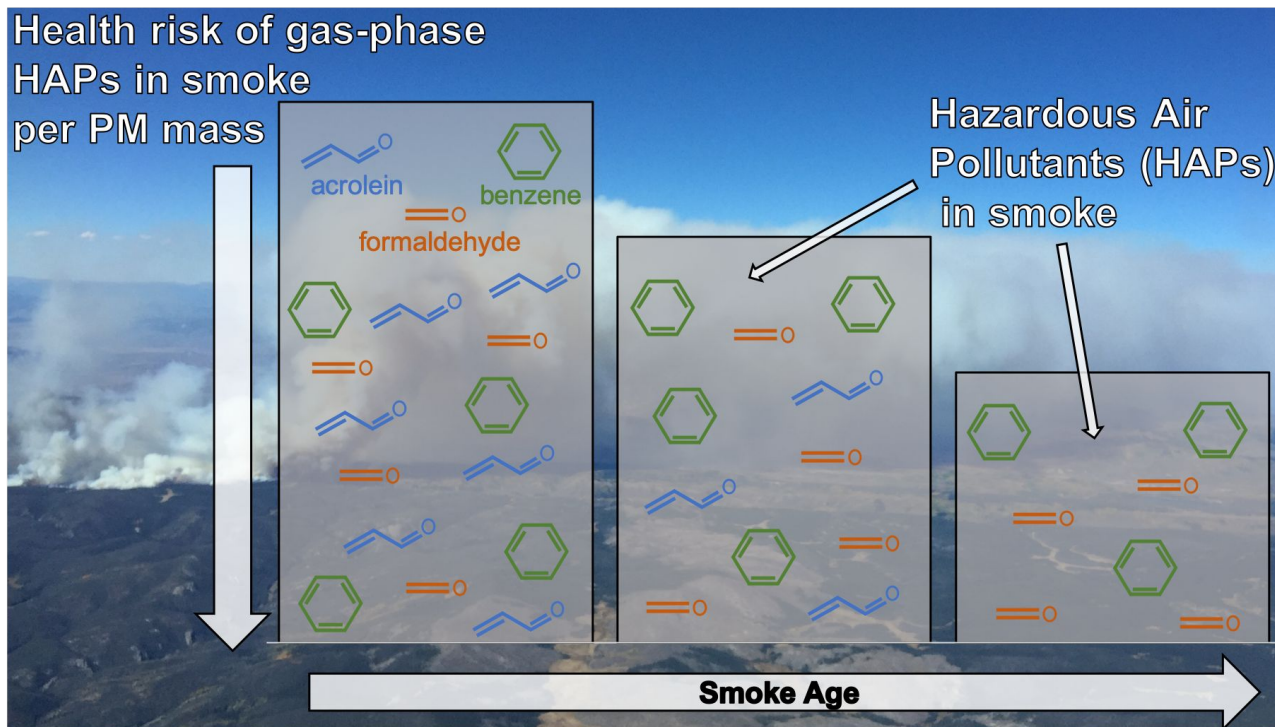


Highly relevant to O_3 in large smoke plumes, we now have a more complete view of daytime NO_y partitioning for the first ~6 hours of evolution.



HONO is the dominant source of OH for the first few hours
Peng et al. (2020)

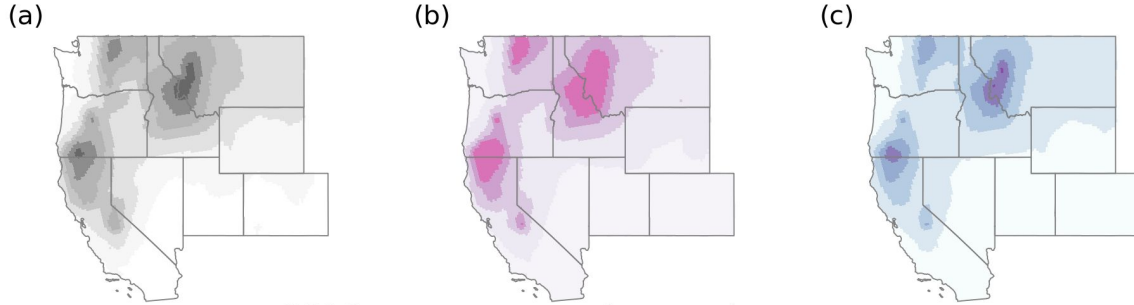
Moving beyond $PM_{2.5}$ and O_3 : Acrolein, formaldehyde, benzene, and HCN are likely the dominant contributors to gas-phase HAPs risk.



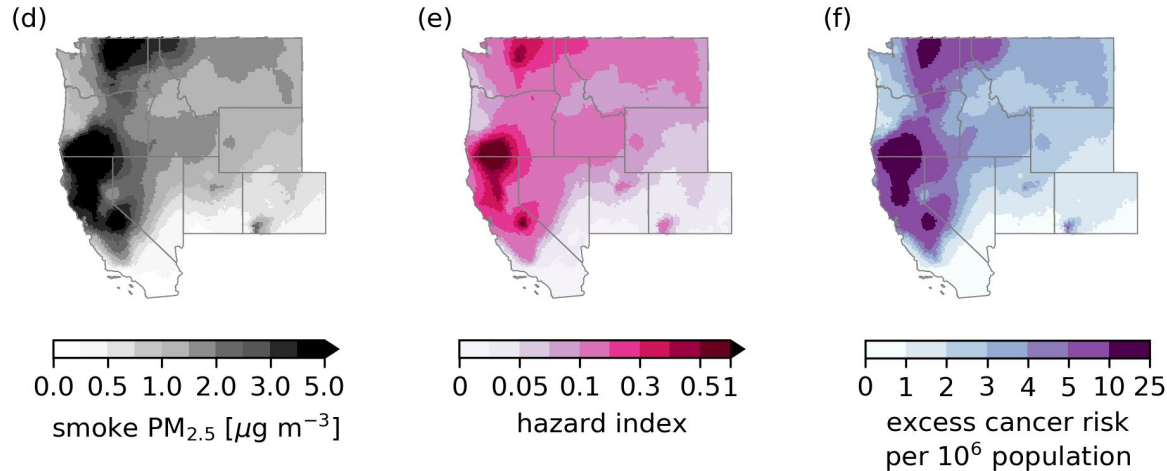
The ratios between acute, chronic noncancer, and chronic cancer HAPs health risk and PM in smoke decrease with smoke age.

A cancer risk of 1 per 10^6 is used to identify locations and pollutants of concern. Heavily fire-prone regions exceed this threshold (mainly due to HCHO).

2006-2018 average smoke exposure

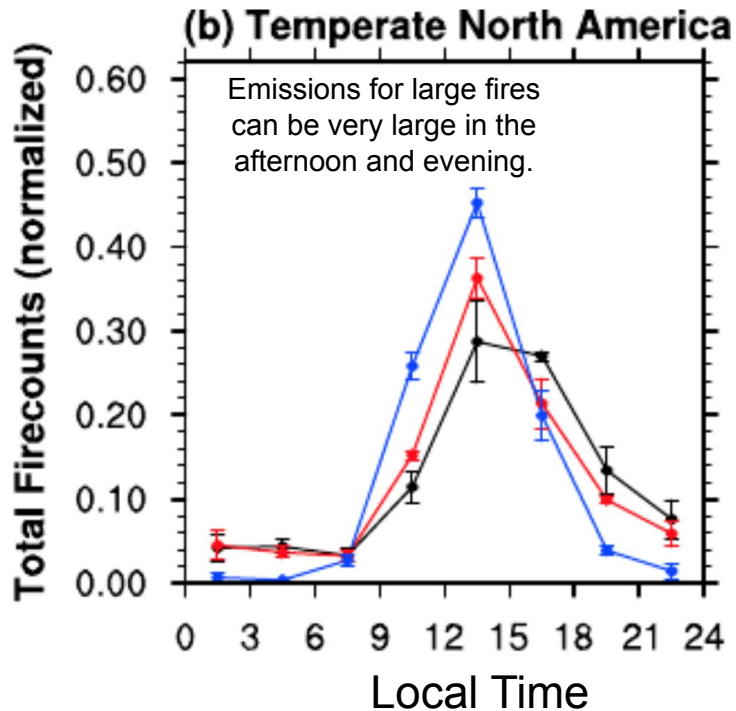


2018 as representative smoke exposure

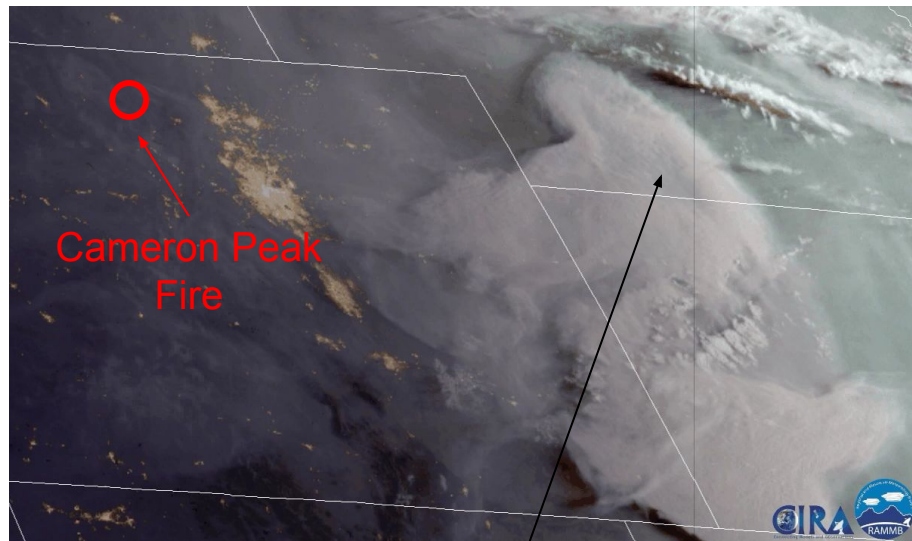


A hazard index of 1 is used as the threshold to indicate a potential health risk. Although smoke-specific hazard indices are below this value, the smoke HAPs may combine with other HAPs sources, or other air pollutants, to increase overall risks.

To understand the full lifecycle of smoke, we also need to consider what happens overnight.



Mu et al. (2011)



Cameron Peak Fire Smoke Plume
Image from 9/7/20
Thanks to Steve Brown and CIRA

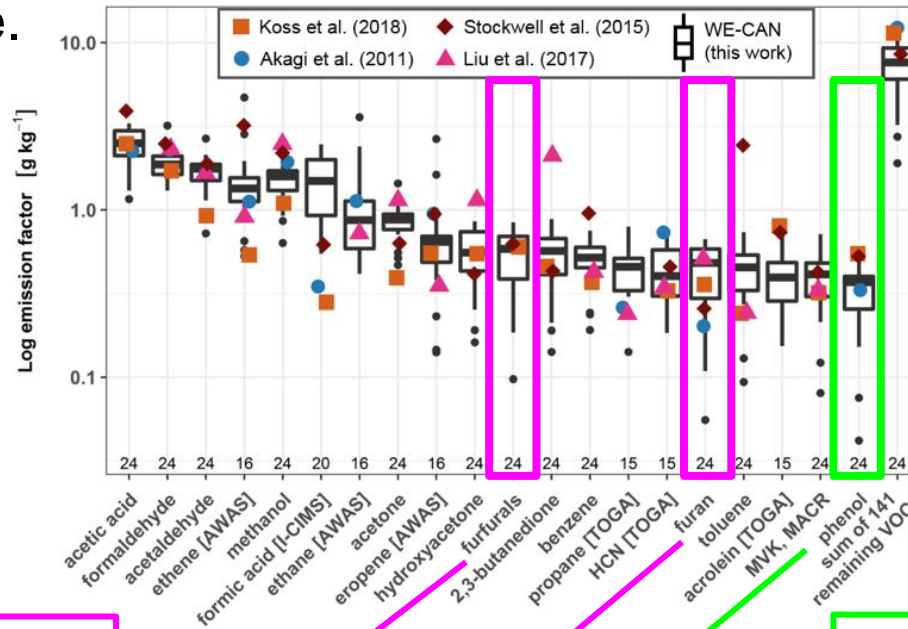
Chemical evolution differs at night when the roles of NO_3 and O_3 can increase.

Furans and furfurals

Oxidized mostly by O_3 or OH
Dominant reactions:

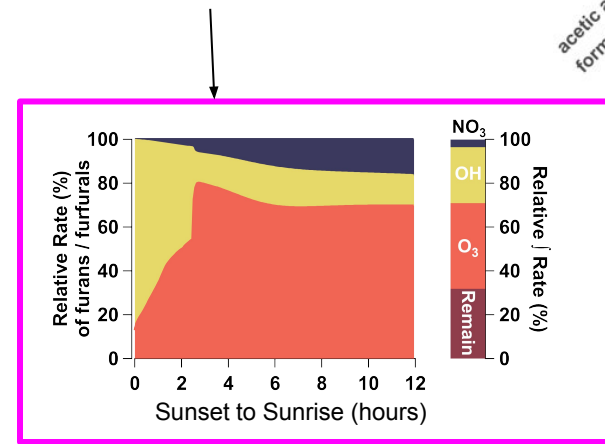
1. $\text{O}_3 + 2,5\text{-dimethylfuran}$
2. $\text{OH} + \text{furanone}$

0 - 30% remain at sunrise

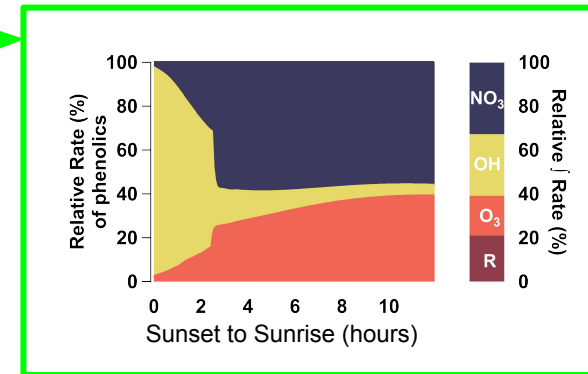


Phenolics

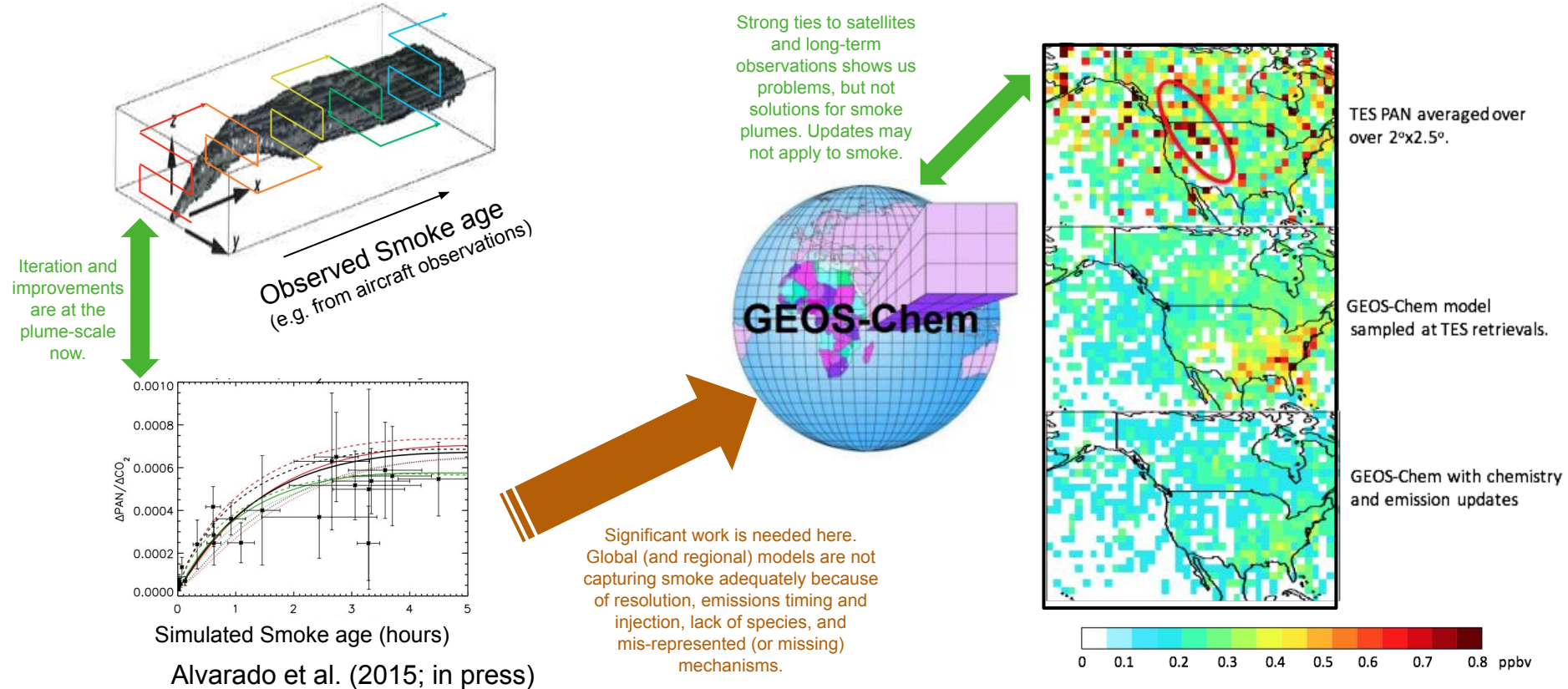
Oxidized mostly by NO_3 or O_3
Catechol and methylcatechol account for 58-77% of phenolic + NO_3 oxidation.
0-20% remain at sunrise



Permar et al. (in review, JGR)
Decker et al. (in prep)



Are our models capturing transformations? Connections between process-level observations and plume-scale models are strong, but links to CTMs are weak.



Summary / Overview Thoughts:

Smoke is a national air quality concern for multiple pollutants.

PM contribution relatively solid, maximum in PNW.

Relevant O₃ chemistry needs work.

Understanding of HAPs is emerging.

Recent major advances in smoke evolution include:

The chemical evolution of OA.

Daytime radical sources

Nighttime chemistry is still a frontier due to under-sampling/study.

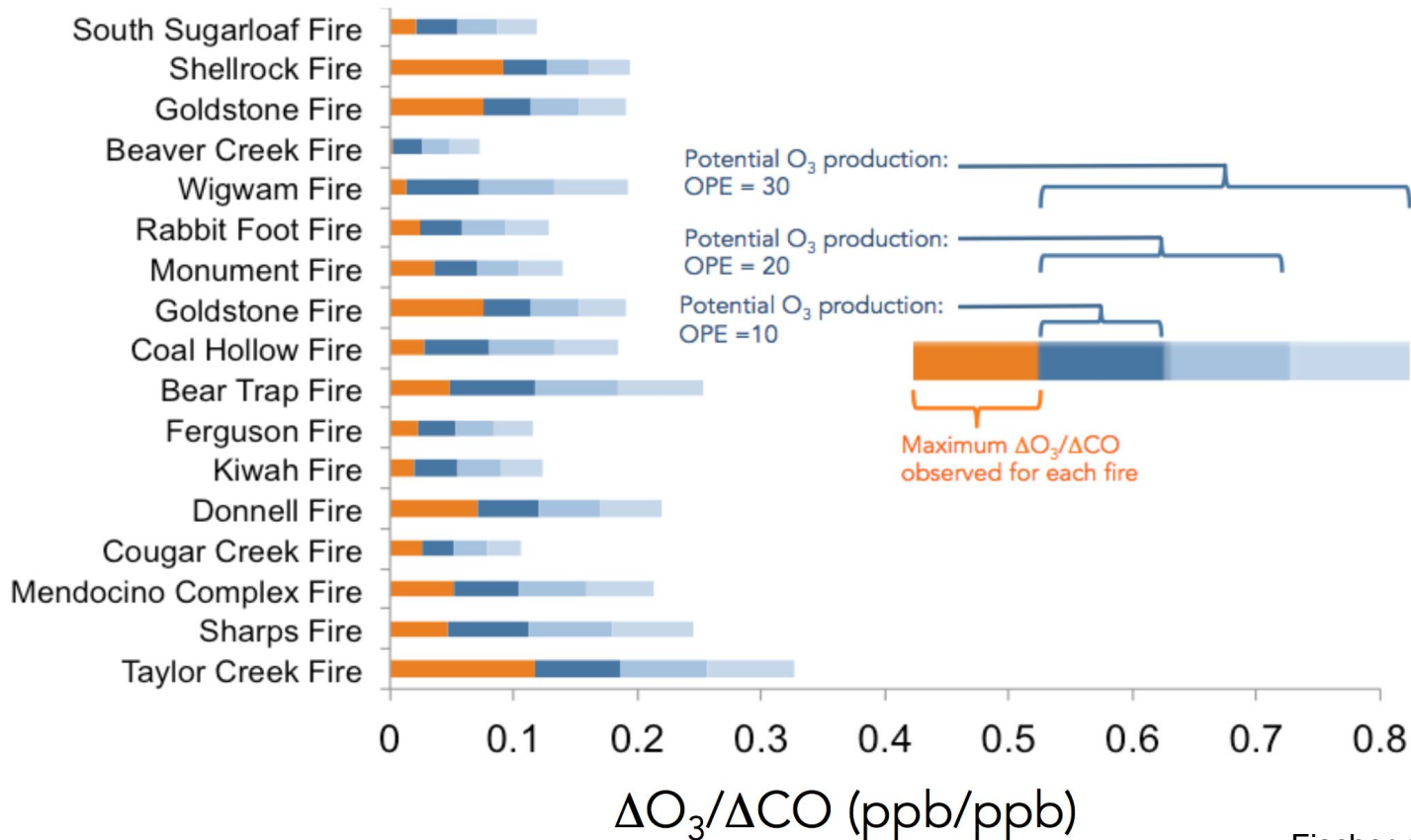
Plume-scale models are improving and can be improved/validated with new data.

Connections to regional/global models are still lacking.

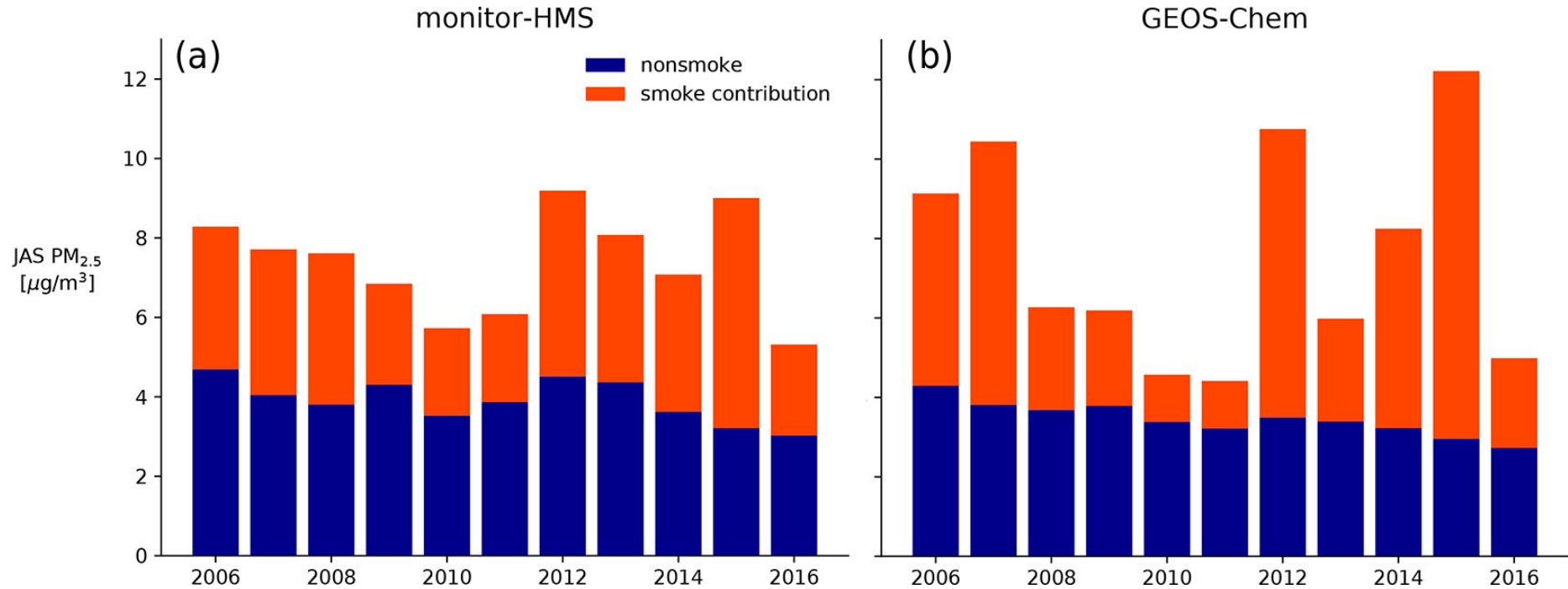
Health-centric simplifications may be possible.

Extra Slides

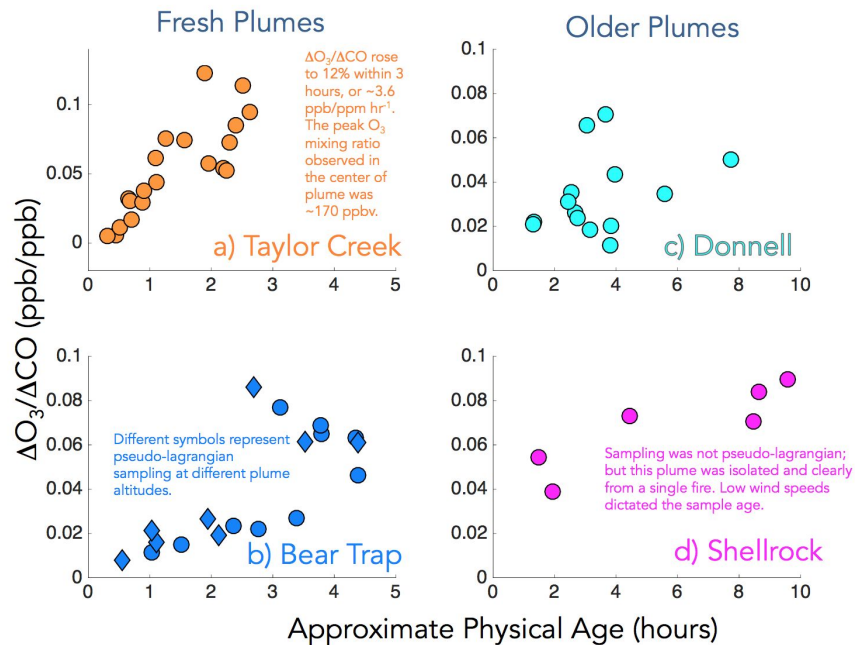
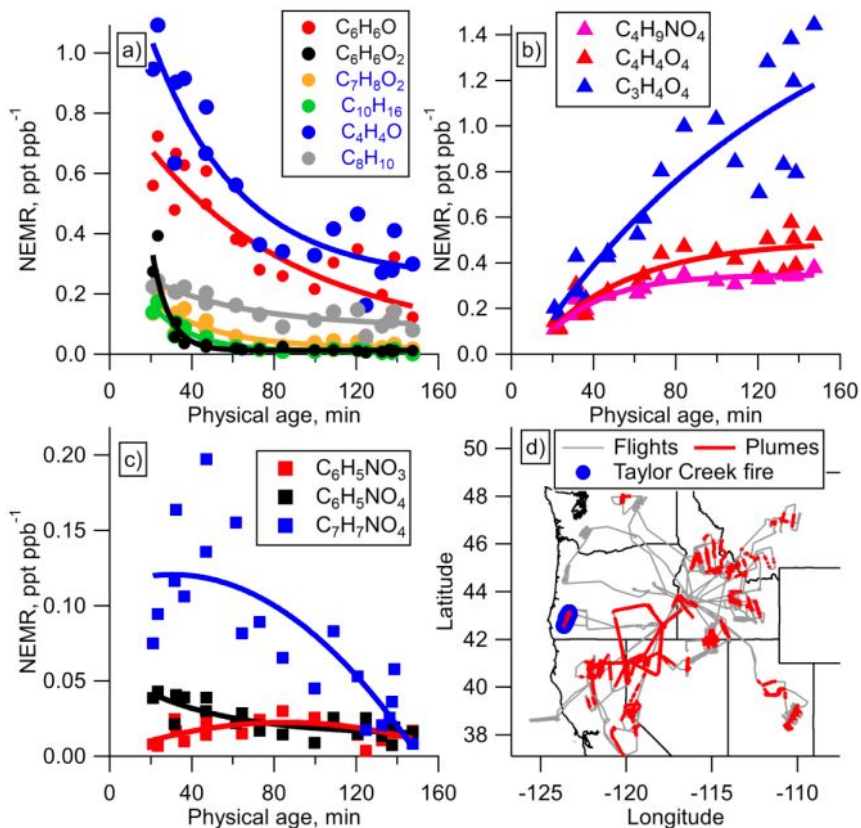
When compared to the potential O₃ from PAN decomposition, the initial bursts of O₃ production likely account for a significant portion of total O₃ production in smoke plumes.



Wildland-fire smoke contributes between ~ 2 and $9 \mu\text{g m}^{-3}$ of the summer-mean $\text{PM}_{2.5}$ in the PNW, depending on the year. In a high-fire year, this is $\sim 70\%$ of the $\text{PM}_{2.5}$.



WE-CAN measurements show rapid (minutes to hours) chemical transformations with implications for OA mass and properties and O_3 formation.



The ratios between acute, chronic noncancer, and chronic cancer HAPs health risk and PM in smoke decrease with smoke age.

