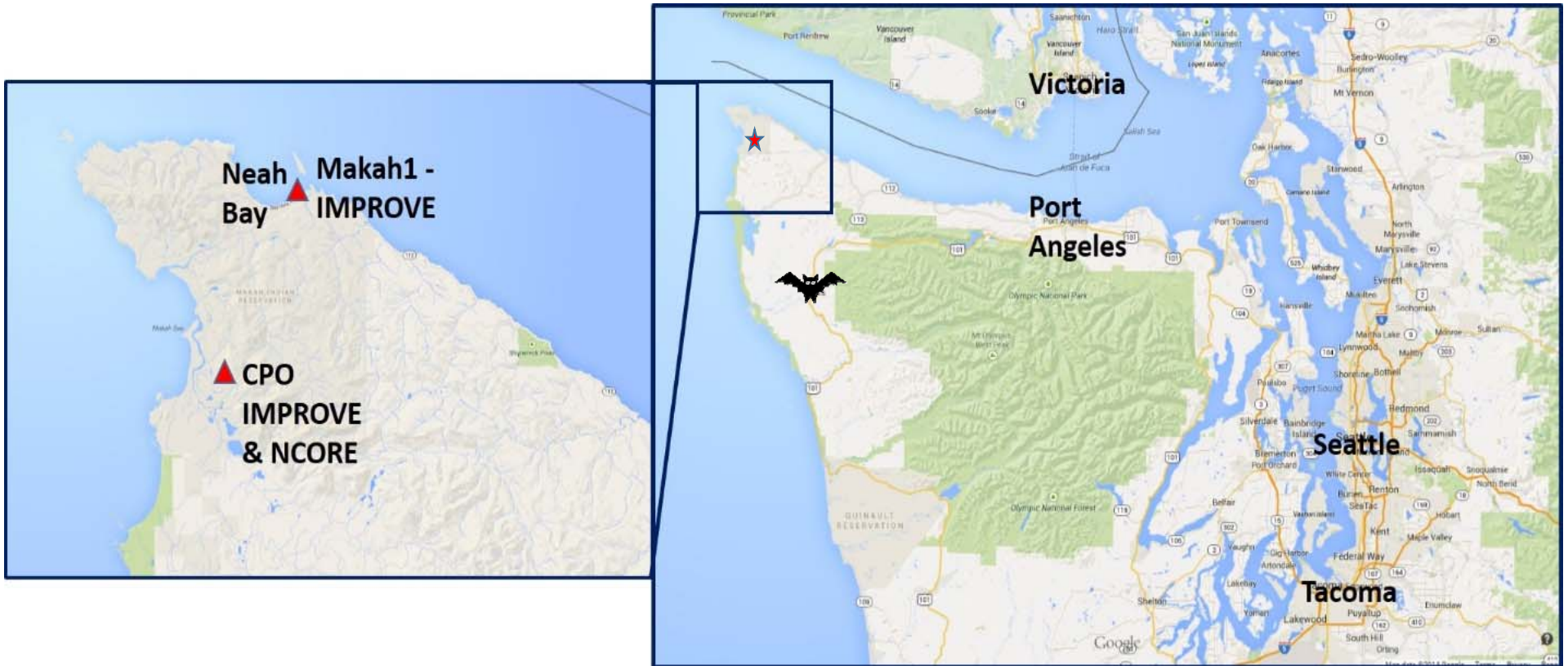
The background image shows a panoramic view from Cheeka Peak. In the foreground, there are dense evergreen trees. The middle ground features a vast expanse of forested hills leading down to a coastline. The ocean is visible in the distance under a clear blue sky. The text is overlaid on this scenic view.

Background PM_{2.5} Source Apportionment at Cheeka Peak Atmospheric Observatory

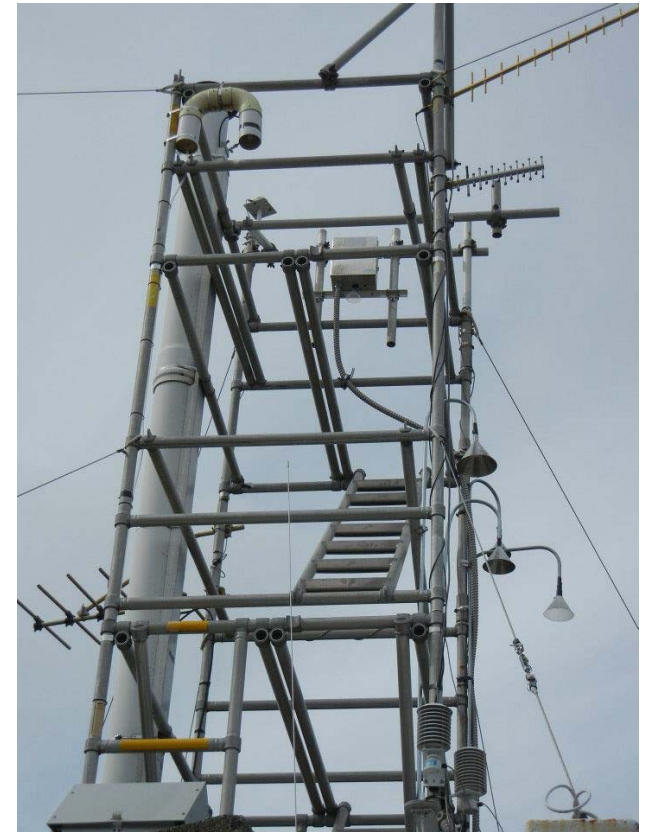
Odelle Hadley, PhD
Senior Air Monitoring Specialist
Olympic Region Clean Air Agency
August 2016

Cheeka Peak Observatory (CPO) Olympic Mountain Range



Project

- Use positive matrix factorization to determine the primary sources of PM_{2.5} at a remote west coast air monitoring site between January 2011 and December 2014
- Compare the resulting source factors with collocated trace gases and meteorological measurements
- Look at long term trends and changes in different source contributions to background PM_{2.5}
- Look at seasonal changes in source contribution



Positive Matrix Factorization (EPA-PMF V5.0)

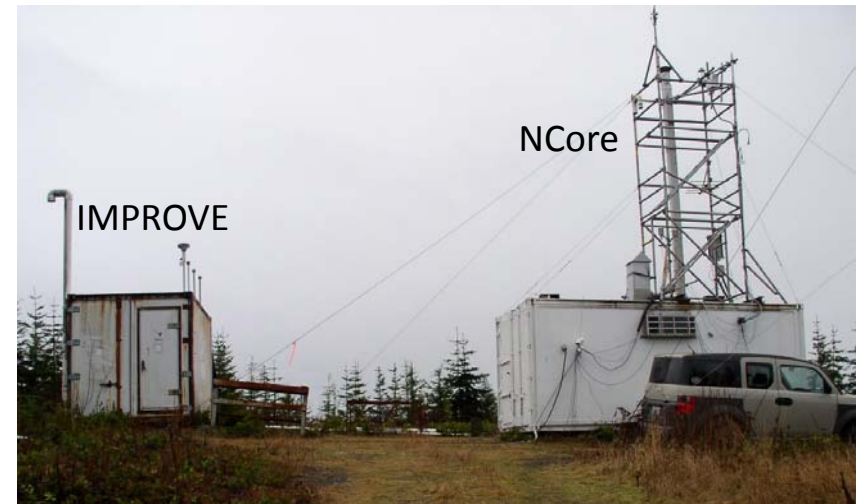
- Receptor based modeling approach
- User determines how many source factors comprise the total aerosol
- Model solves equation 1 iteratively (30 runs) to minimize Q (eqn 2):

$$1) x_{ij} = \sum_{k=1}^n g_{ik} f_{kj} + e_{ij} \qquad 2) Q = \sum_{i=1}^n \sum_{j=1}^m \left[\frac{x_{ij} - \sum_{k=1}^p g_{ik} f_{kj}}{u_{ij}} \right]^2$$

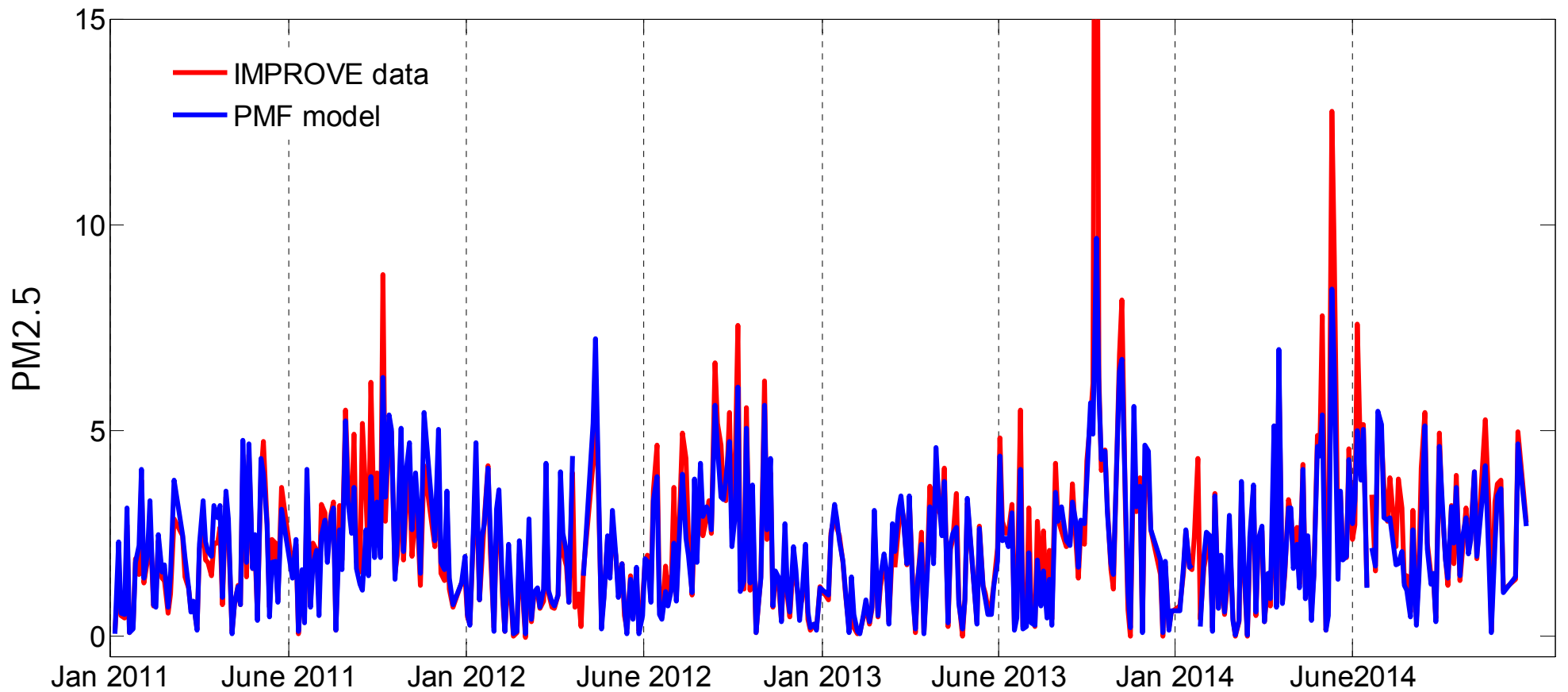
- x_{ij} is the input data, where i is the number of days for which a sample was collected and j is the number of chemical species in a sample
- g is the mass each factor (k) contributes to each sample
- f is the fraction each chemical species contributes to a source factor
- e_{ij} is the residual of each species for a given day
- PM2.5 is the “total mass variable”
 - Forces the sum of the source factors to equal the measured PM2.5 input data

Data collected at CPO

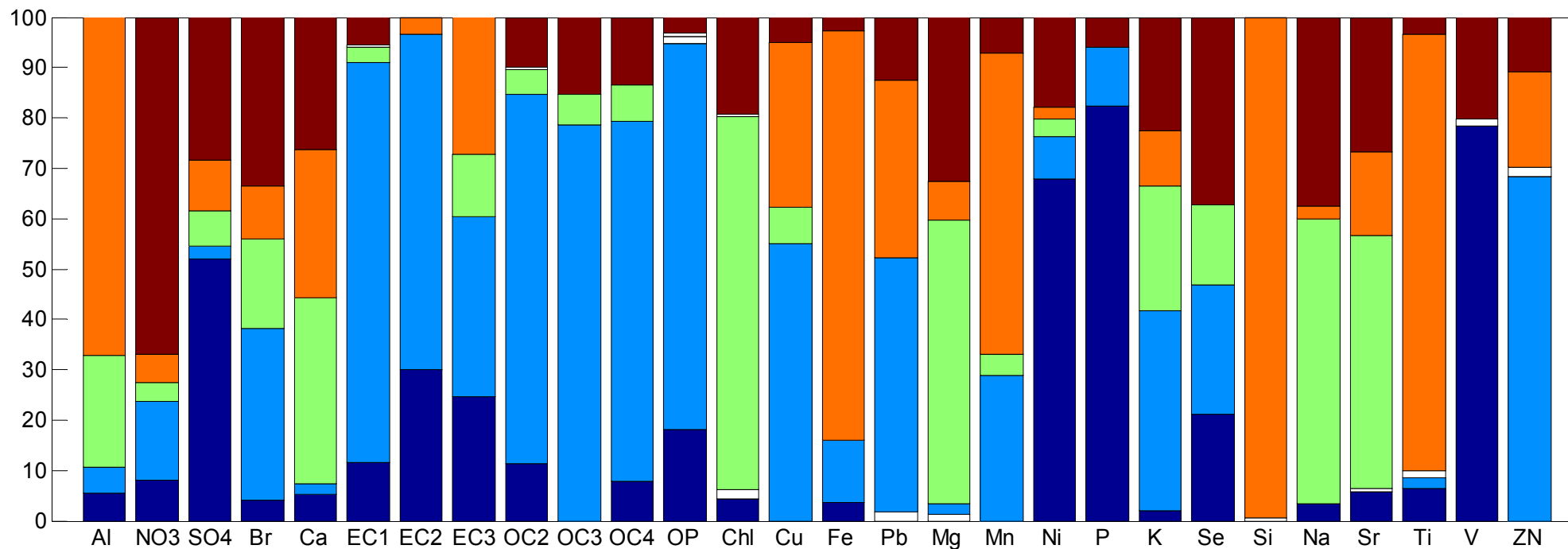
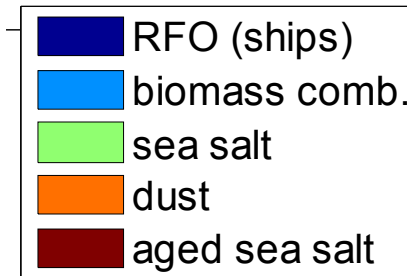
- IMPROVE data are from PM2.5 filters representing a 24-hour average of chemically speciated particulate collected once every three days
 - 28 chemical species (excluding PM2.5)
 - Was used as input to the PMF model
- NCORE data - hourly resolved trace gas, meteorological, and light scatter data
 - **CO, SO₂, NO/NO_y, & O₃**; nephelometer derived PM2.5; **wind speed and direction**, temperature, pressure, & RH
 - These data helped place PMF model results into context



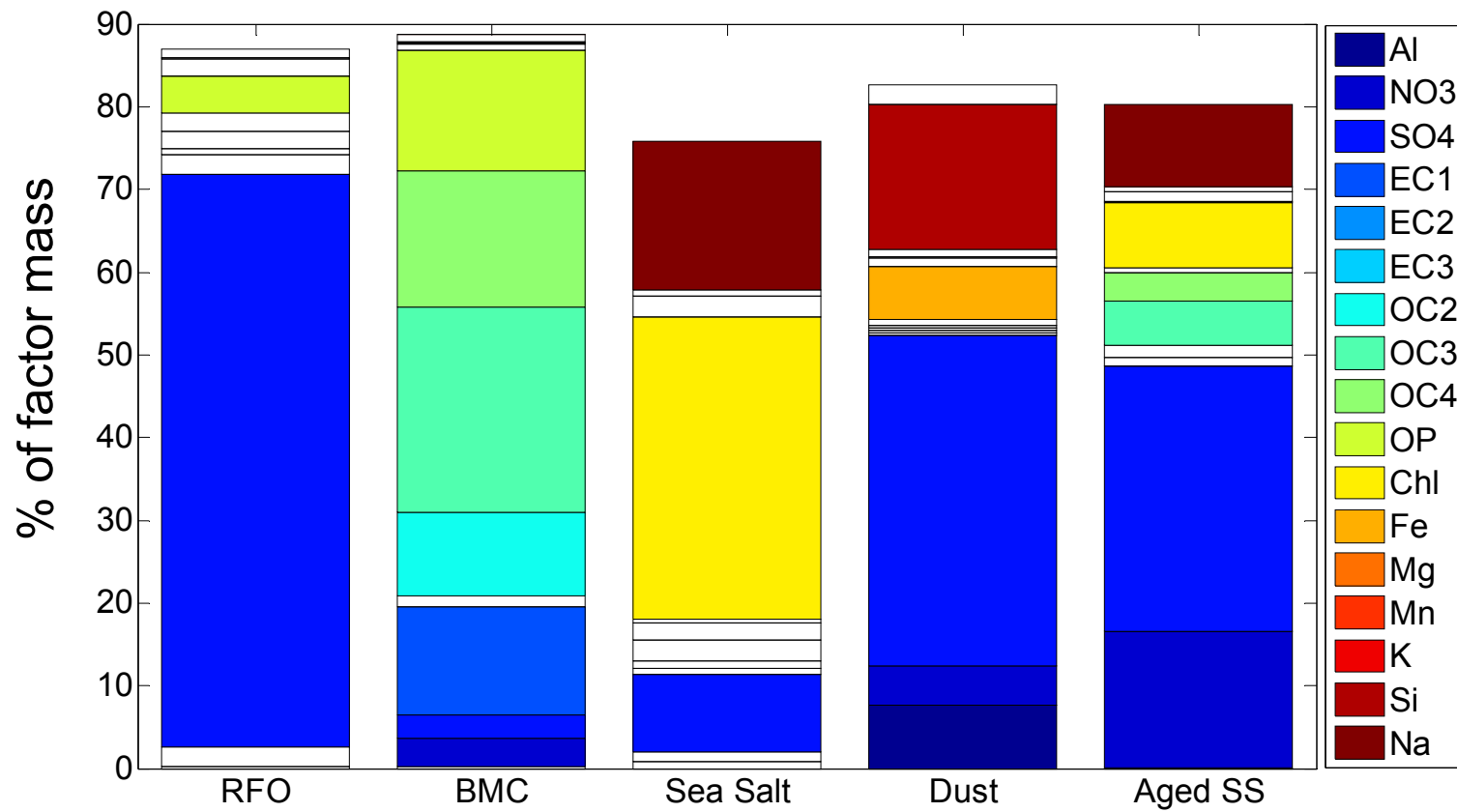
Measured vs. modeled PM2.5



Chemical Fingerprints of Source Factors



Primary components in each source factor

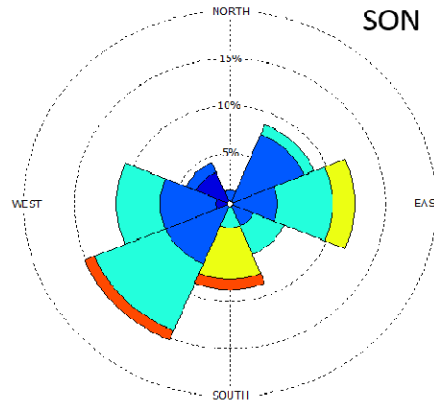
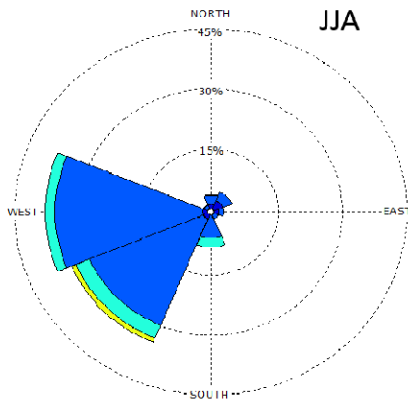
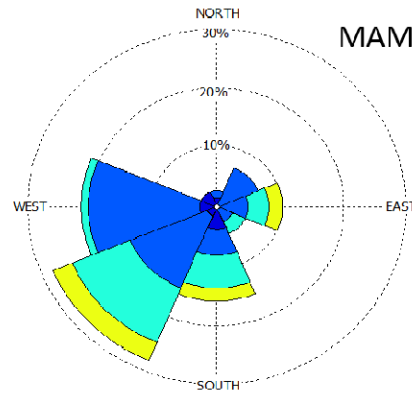
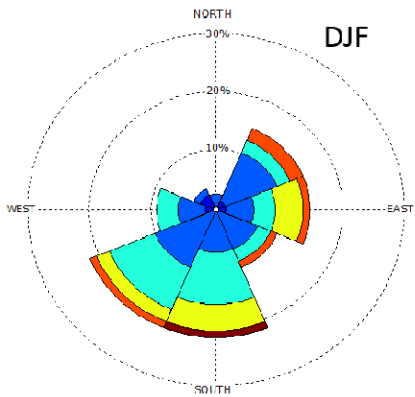


Error Estimation

- Displacement error estimation
- DISP Diagnostics:
 - Error Code: 0
 - Largest Decrease in Q: -0.105
 - %dQ: -0.000423
 - Swaps by Factor: 0 0 0 0 0 (dQmax = 4, 8, 15, 25)
- Bootstrap error estimation (R=0.7 [0.6]), 100 bootstraps

	RFO_base	BMC_base	SS_base	Dust_Base	Aged_base	unmapped
Boot -RFO	71 [98]	8[0]			4 [2]	17 [0]
Boot -BMC		100				
Boot -SS			100			
Boot -Dust				100		
Boot -Aged					100	

Seasonal wind rose at Cheeka Peak



Wind Speed
(MPH)

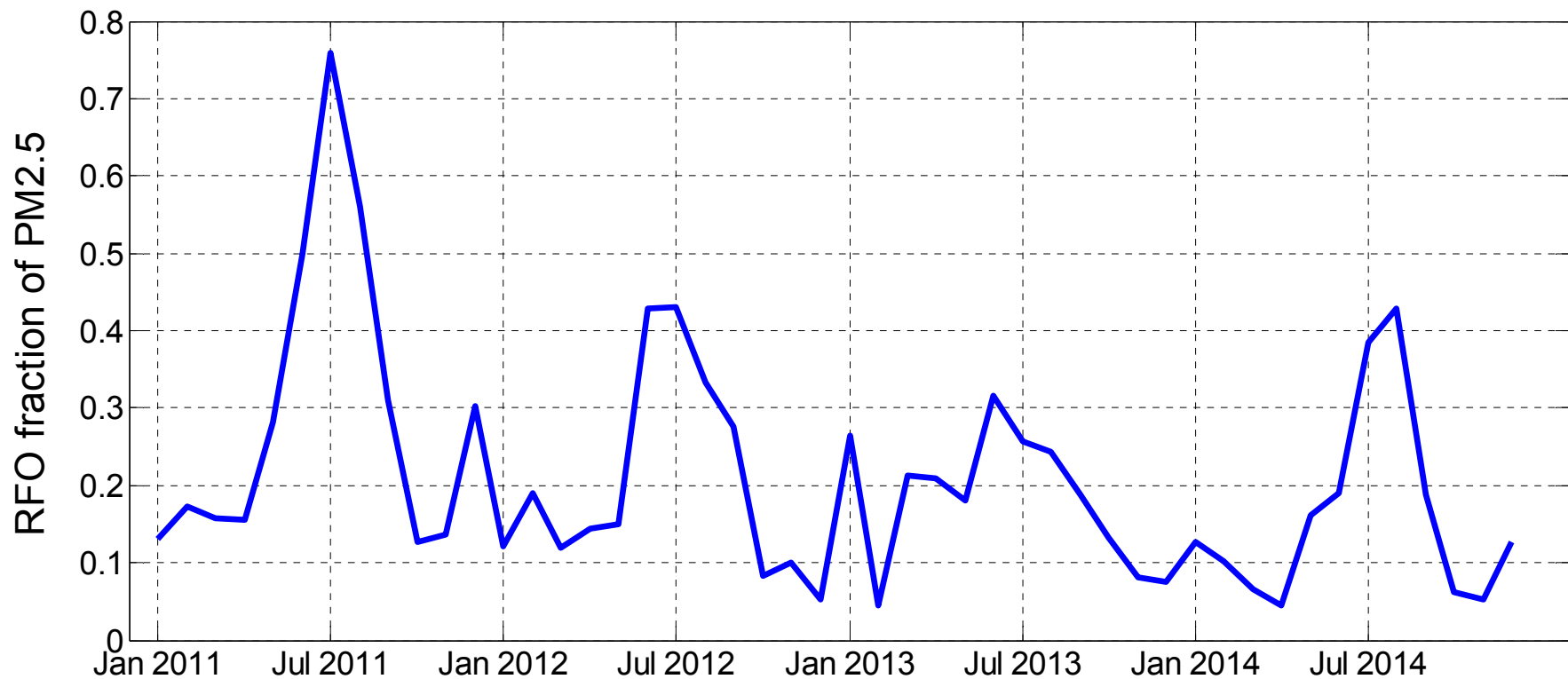


- Westerly winds dominate
- Easterly winds are most frequent in winter and fall
- Summer winds are almost exclusively westerly or southwesterly
- Highest wind speeds in winter and fall

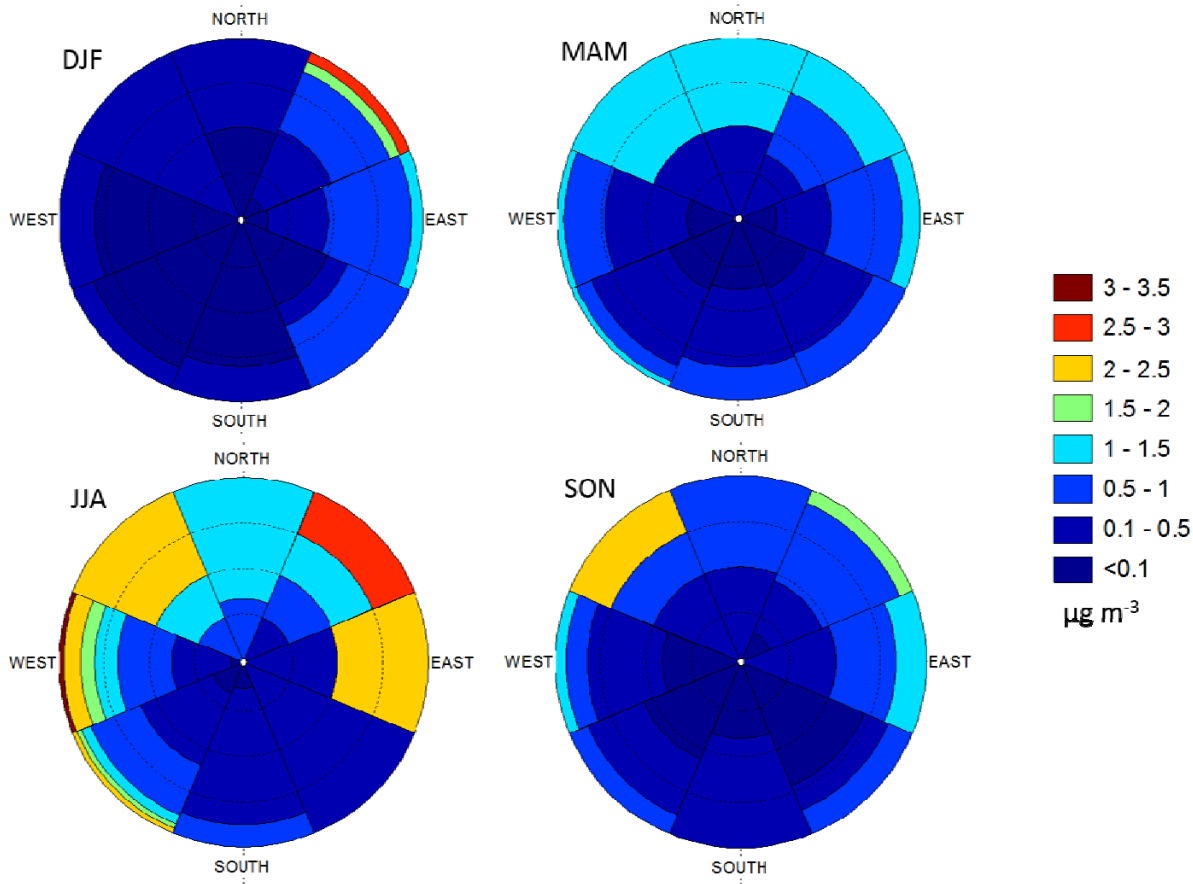
Long term and seasonal trends Residual fuel oil (RFO) a.k.a marine vessel emissions



Long term and seasonal trends Residual fuel oil (RFO) a.k.a marine vessel emissions

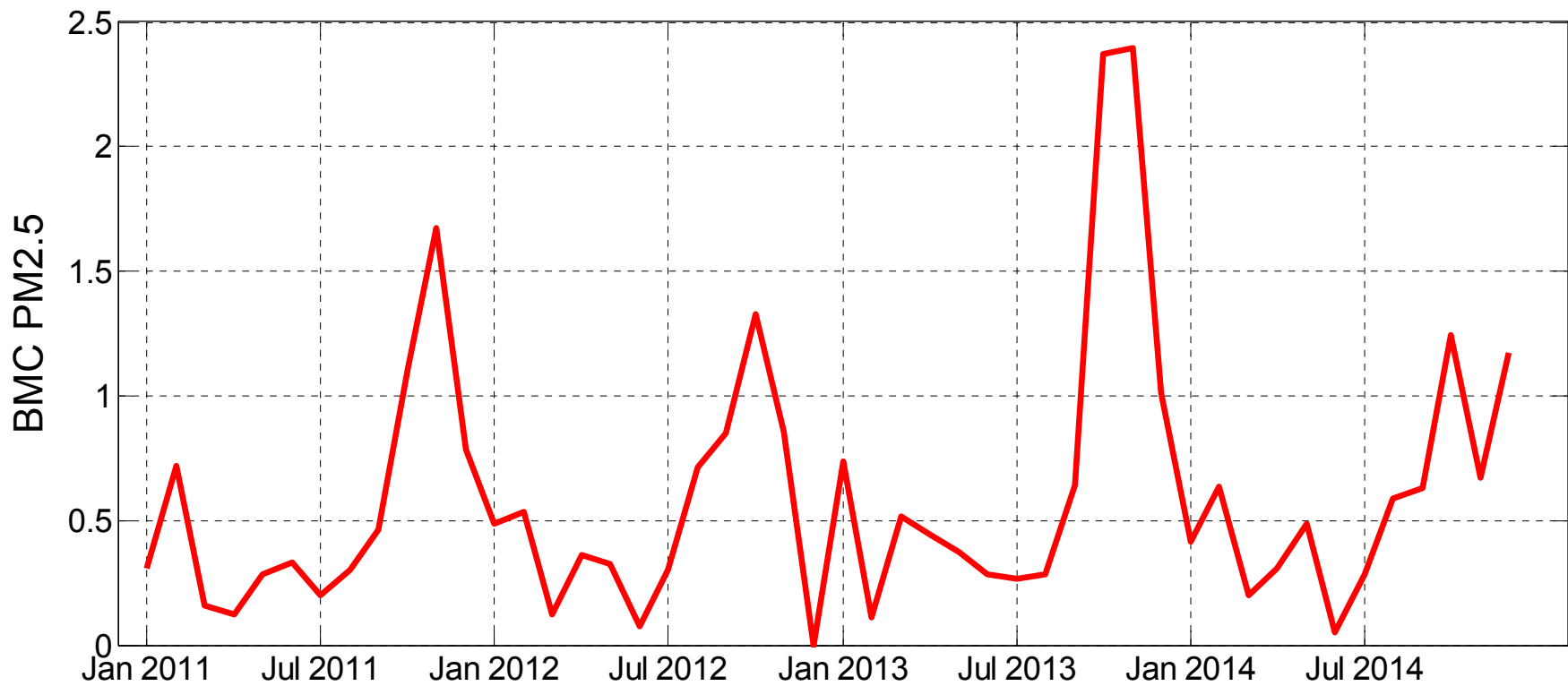


Seasonal wind direction - RFO



- Normalized frequency distribution with respect to wind direction
- Highest concentrations follow shipping routes:
 - West of CPO
 - North through Strait of Juan de Fuca
 - East and Northeast ports of Tacoma, Seattle, and Vancouver
- Highest concentrations observed in the summer

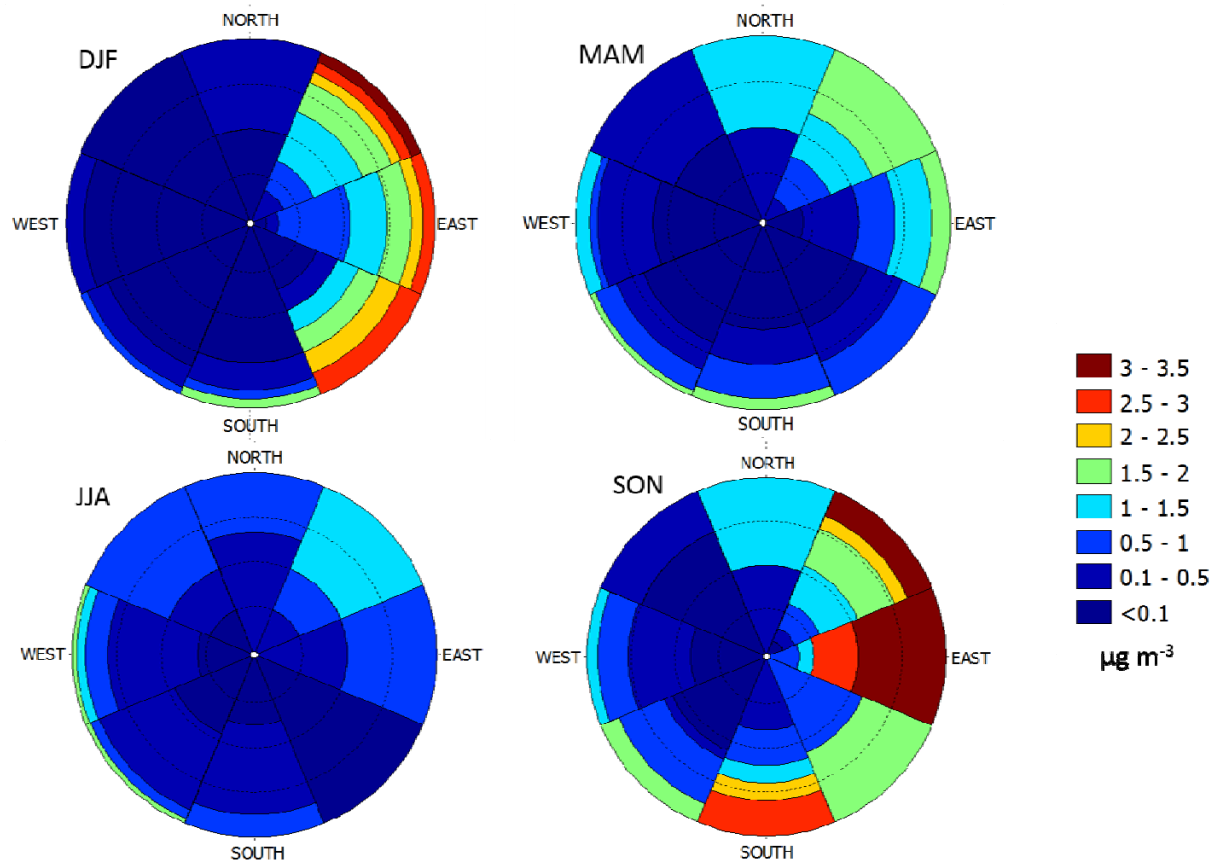
Long term and seasonal trends biomass combustion (BMC)



Long term and seasonal trends biomass combustion (BMC)

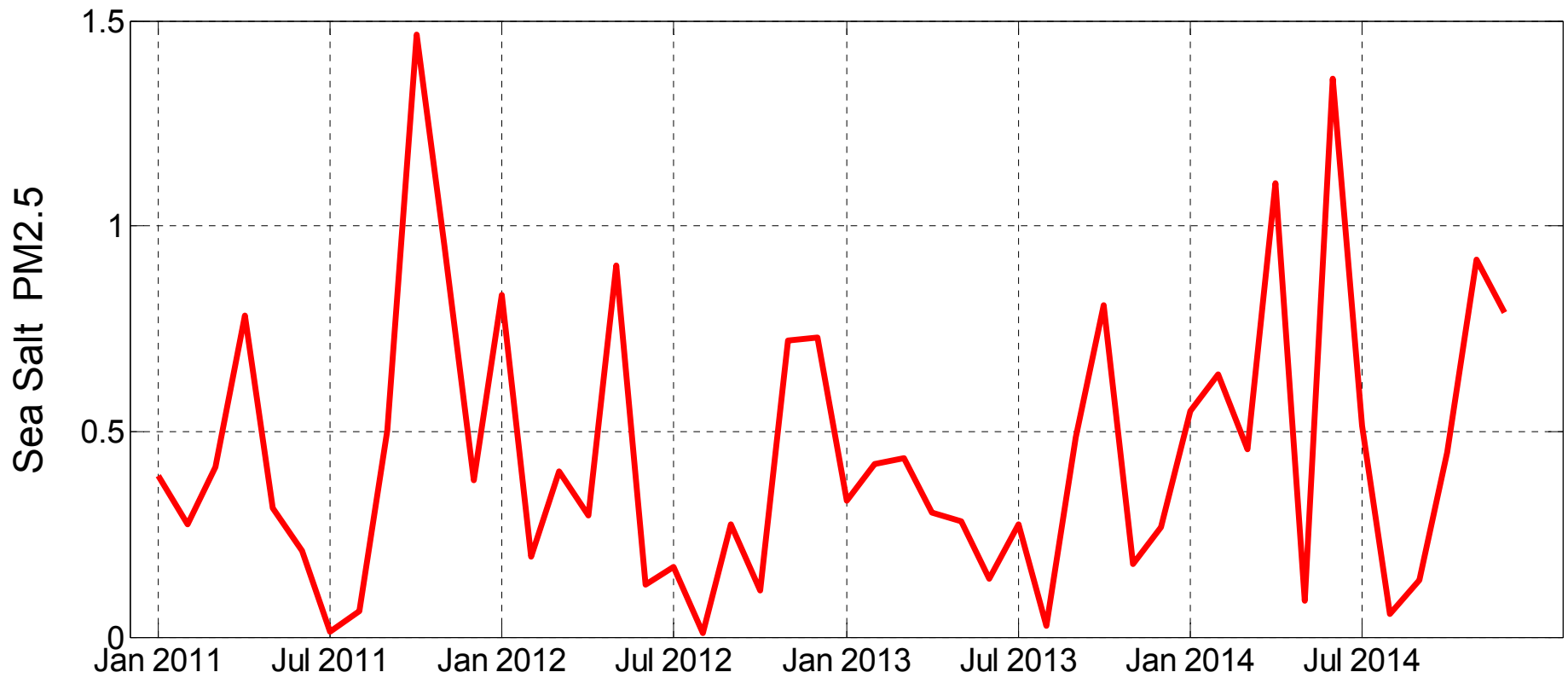


Seasonal wind direction - BMC

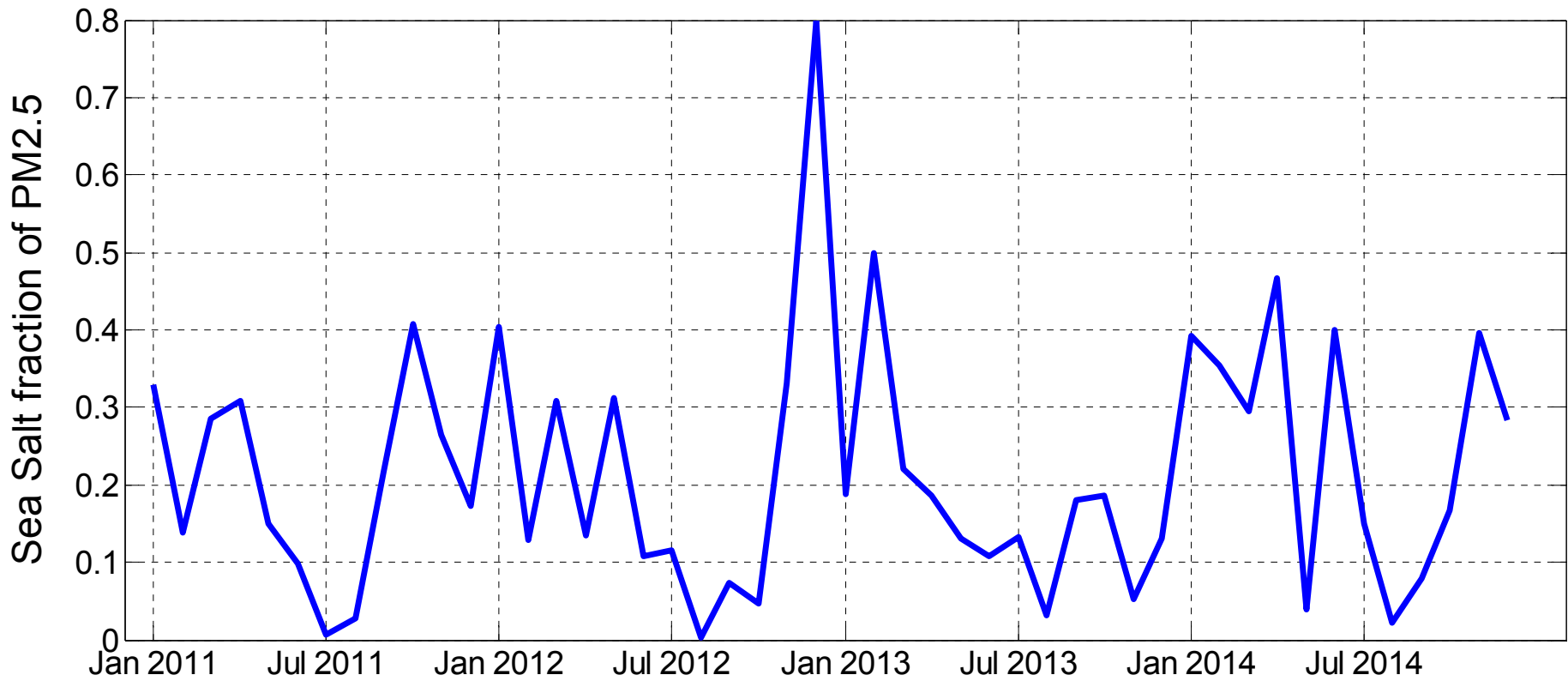


- Highest concentrations are associated with easterly winds:
 - Continental air
- Highest concentrations observed in winter and fall
 - Winter concentrations likely associated with residential burning
 - Fall concentrations with outdoor burning and residential heating

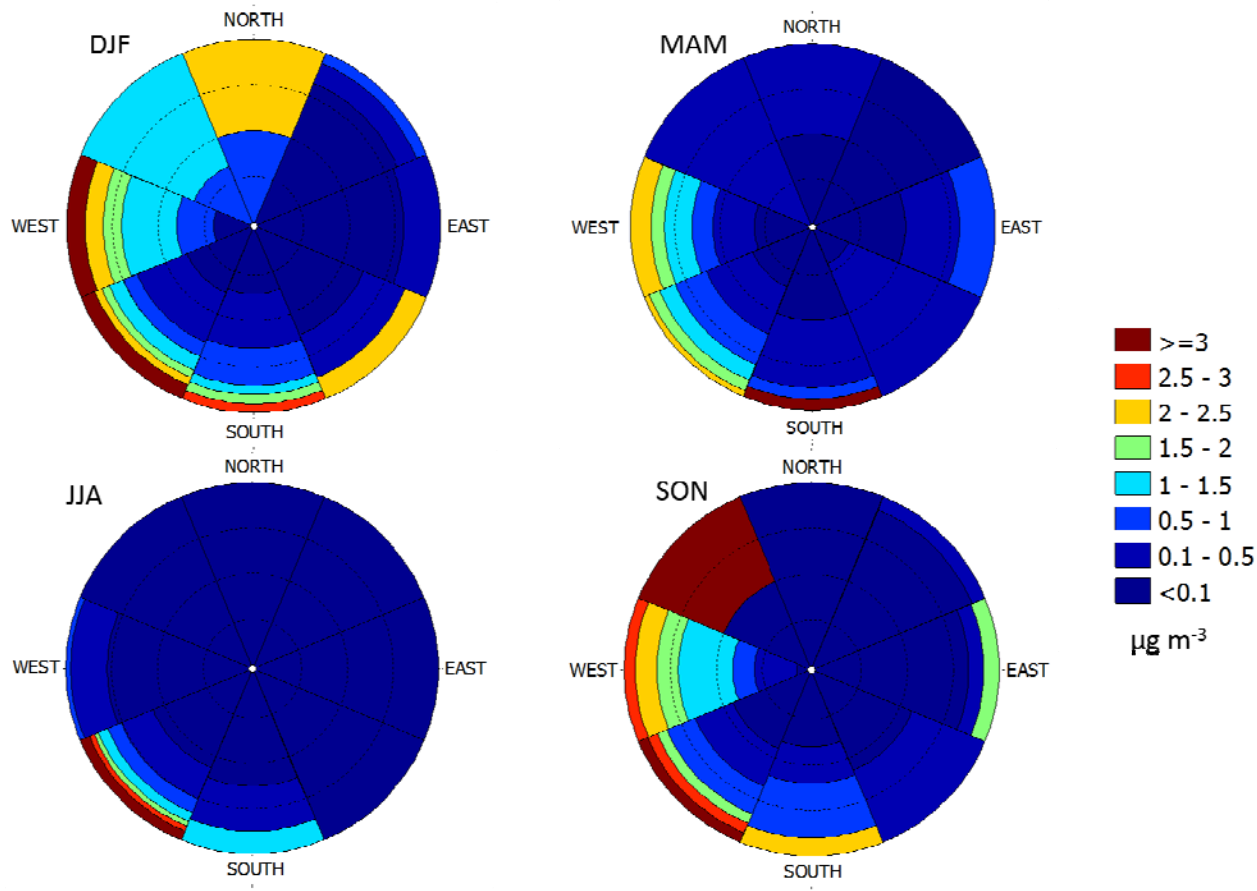
Long term and seasonal trends – Sea Salt



Long term and seasonal trends – Sea Salt

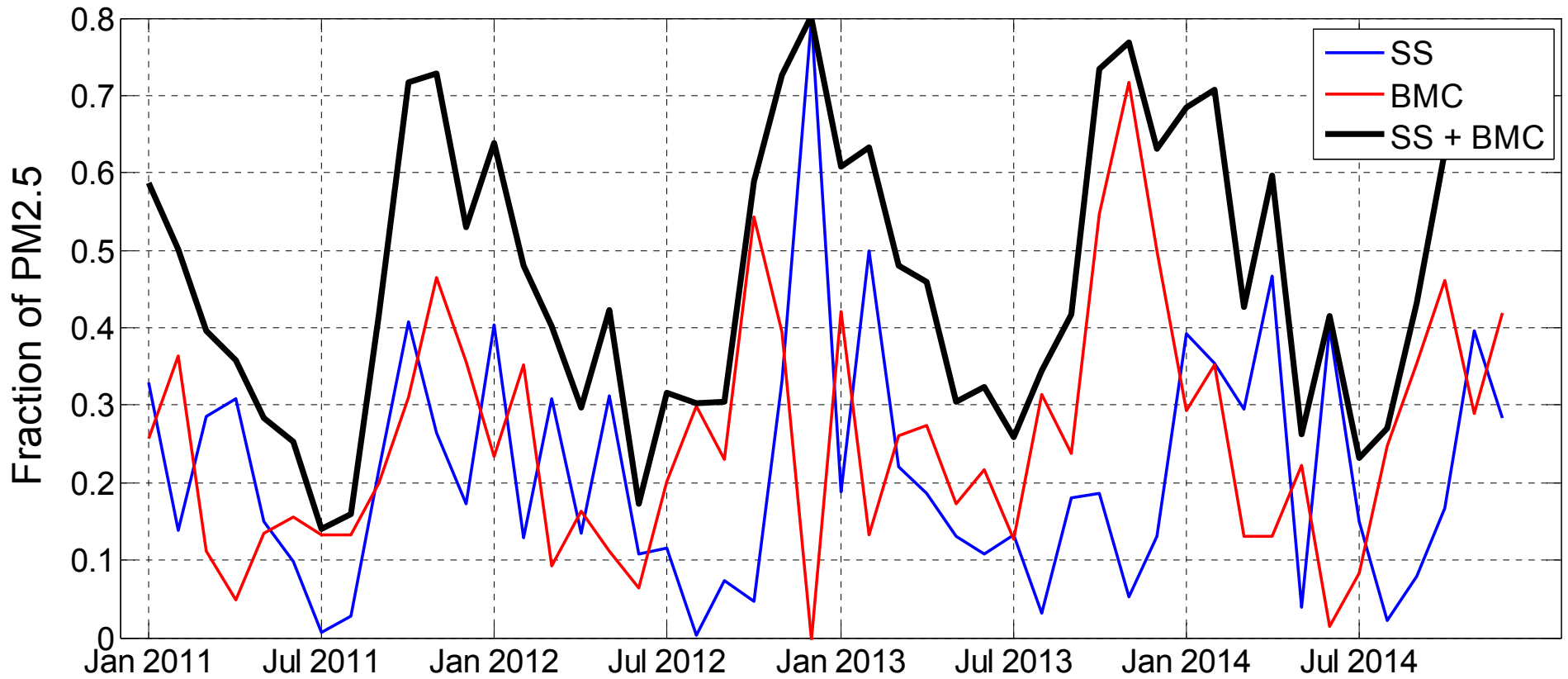


Seasonal wind direction – Sea Salt

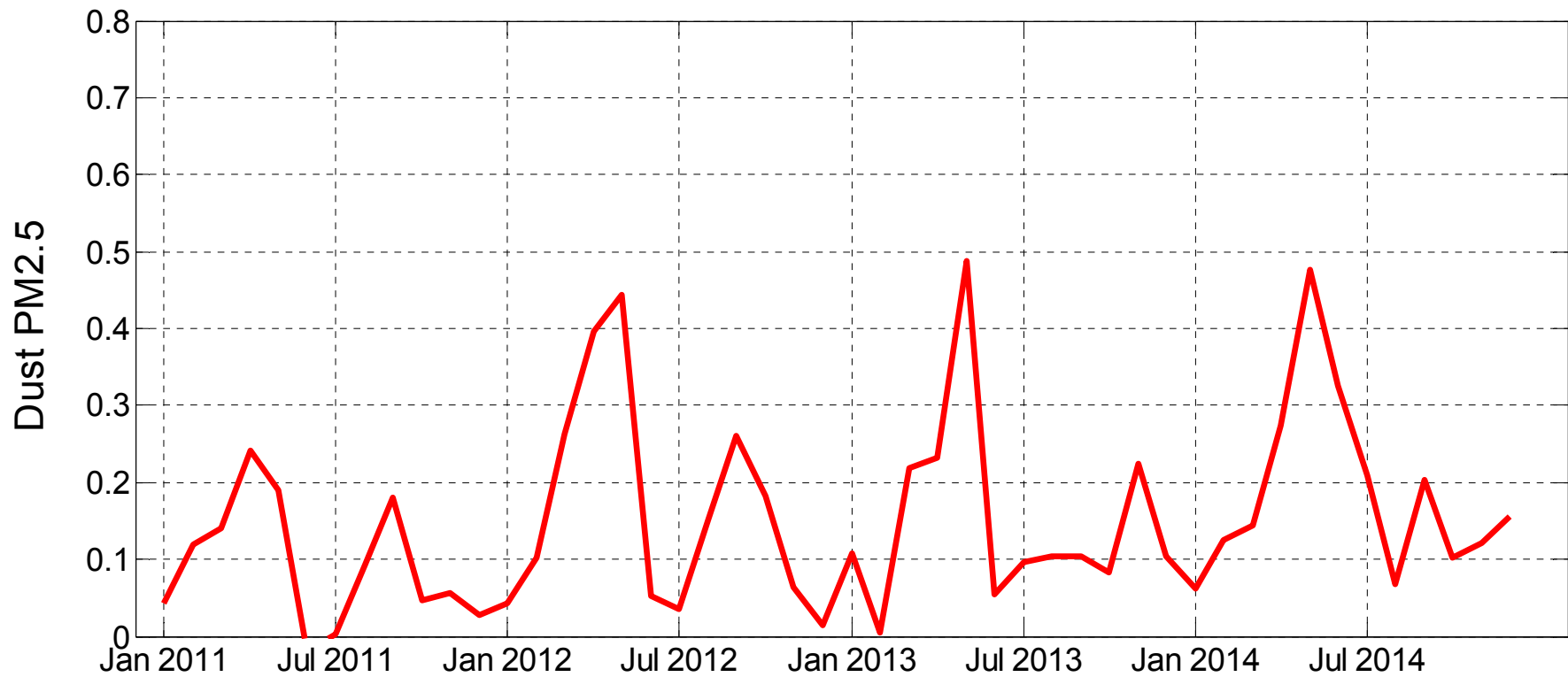


- Highest concentrations are associated with westerly winds:
 - Marine air
- Highest concentrations observed in winter and fall
 - Increase in marine storms (wind speed & wave breaking)

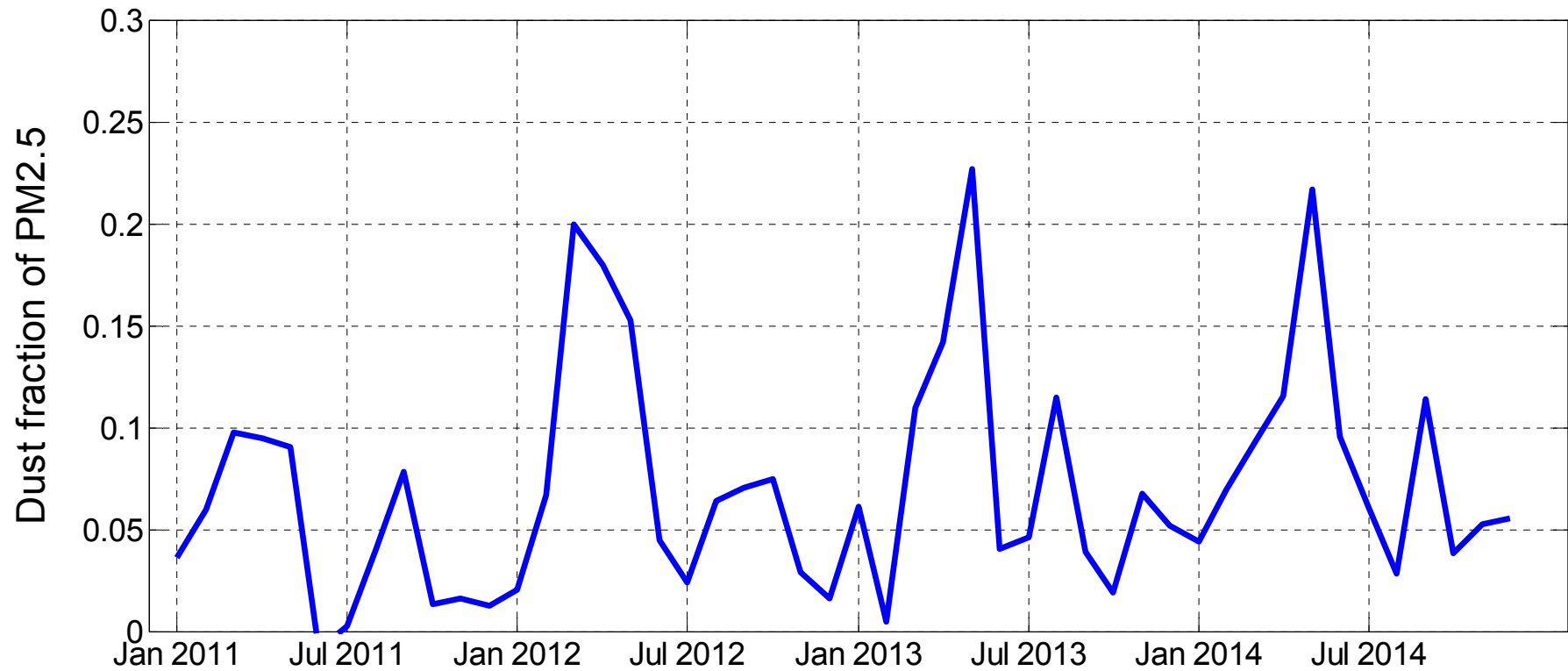
Sea Salt and BMC are the primary fall/winter pollutants



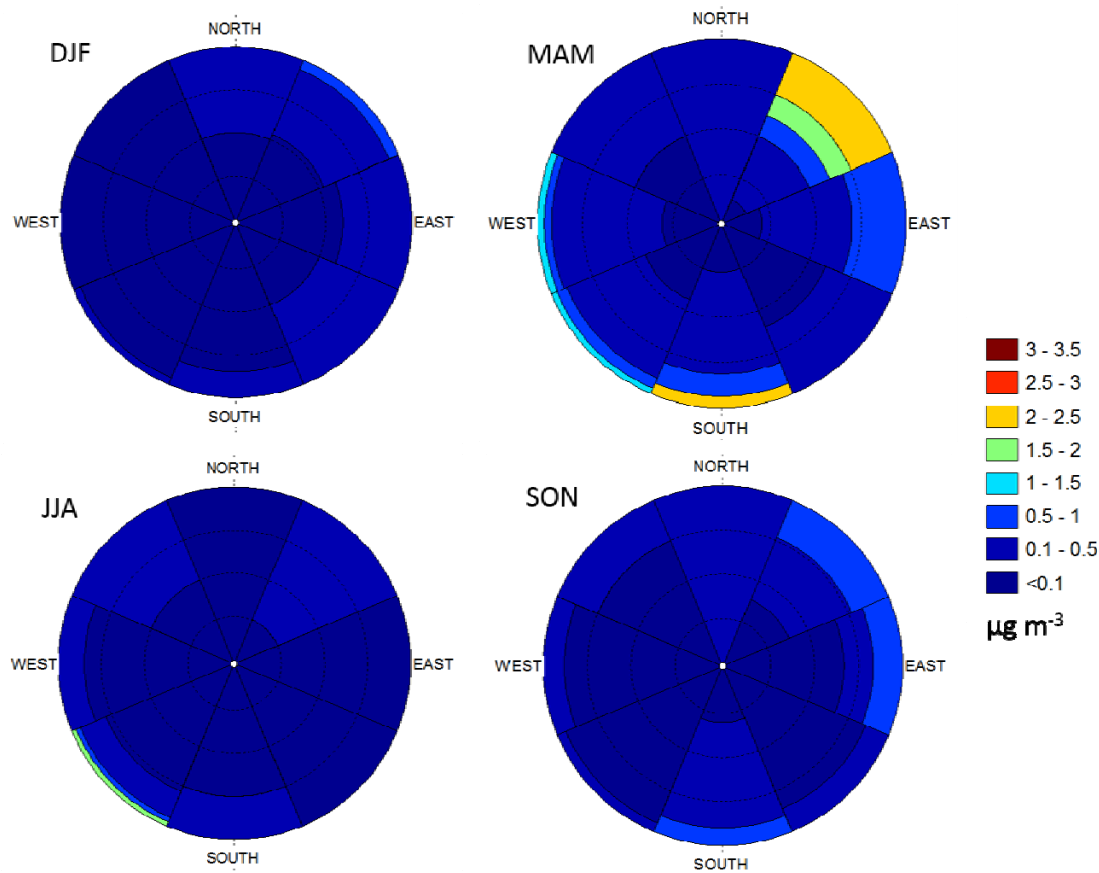
Long term and seasonal trends – Dust



Long term and seasonal trends – Dust

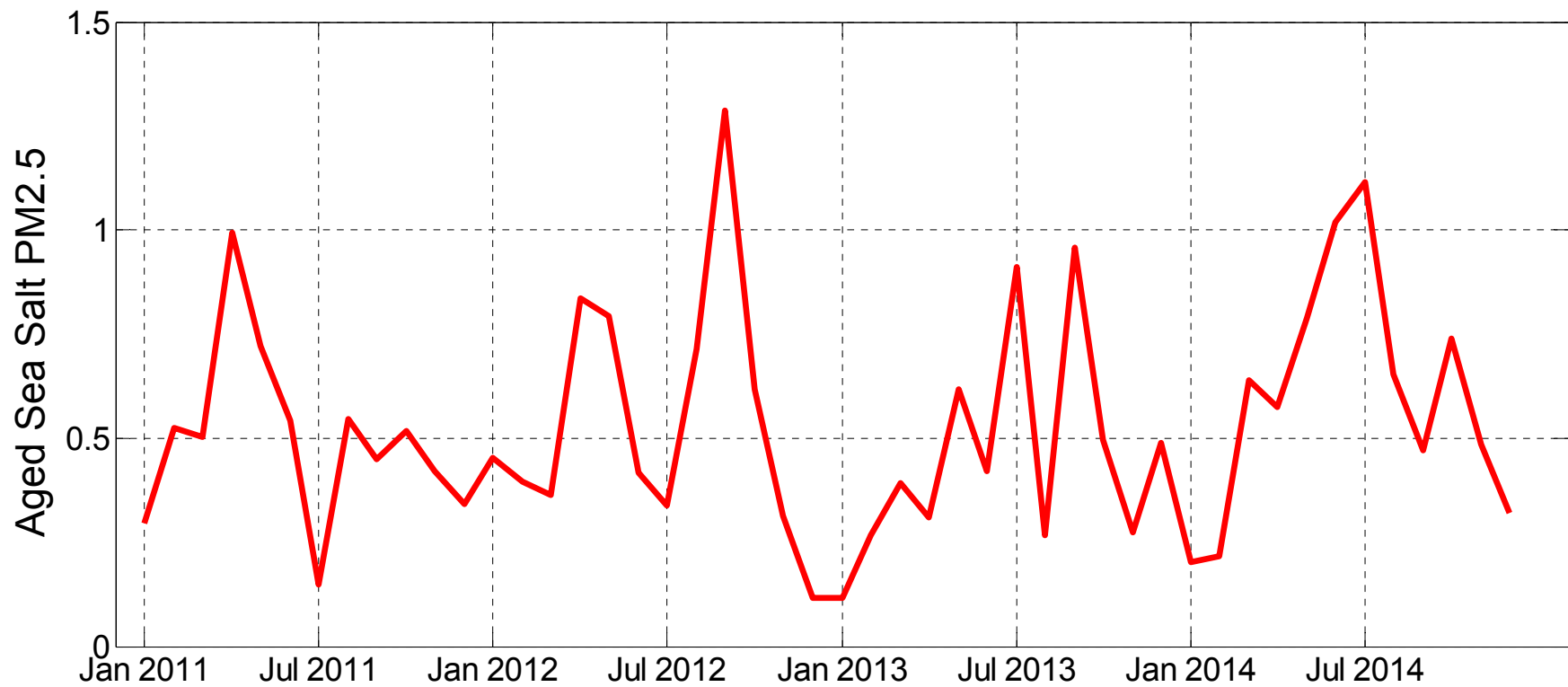


Seasonal wind direction – Dust

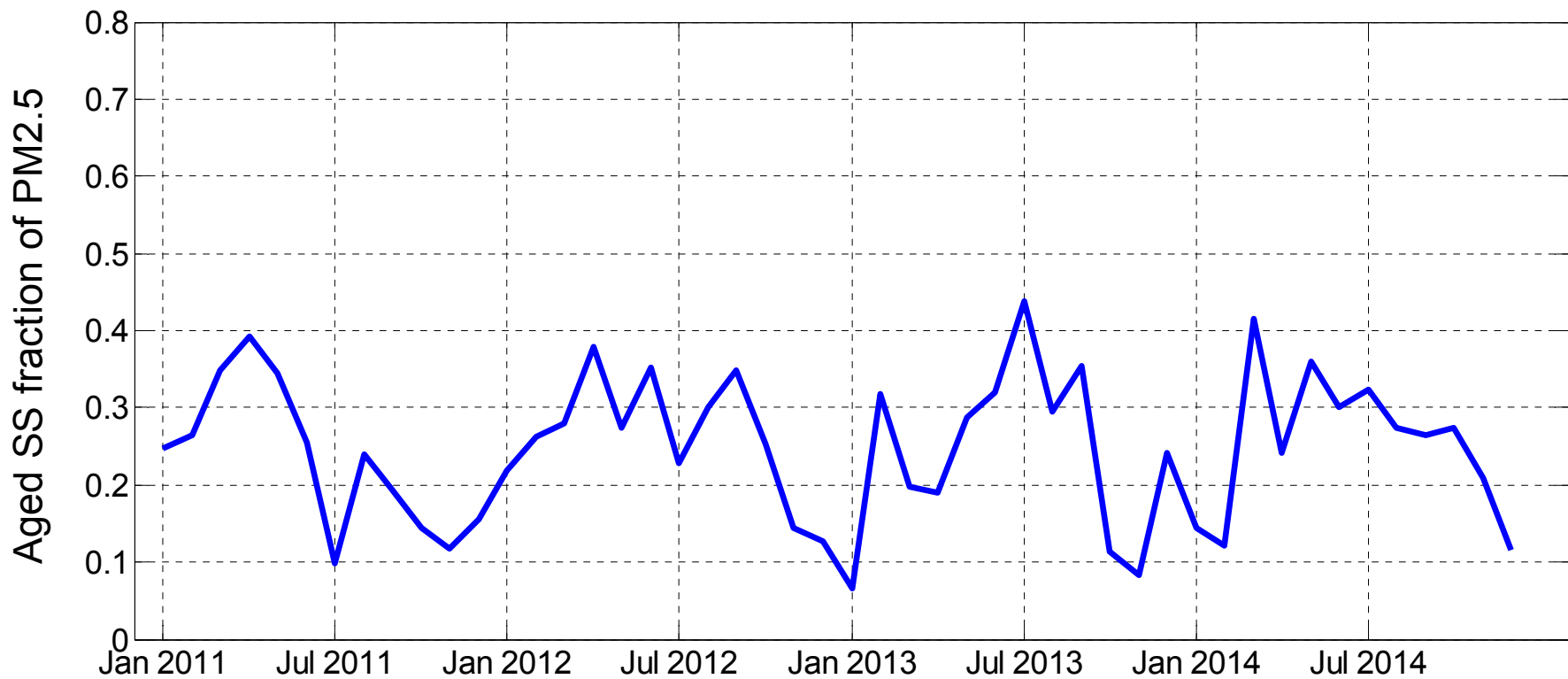


- Highest concentrations are associated with NE winds
- Highest concentrations observed in spring
 - Coincides with spring trans-Pacific transport (TPT) season
 - Surface wind direction is inconsistent with TPT (*Hadley et al 2007*)
 - Fe/Ca exceeds 2 indicating North American source (*VanCuren et al. 2005*)
 - Trajectory analysis is needed

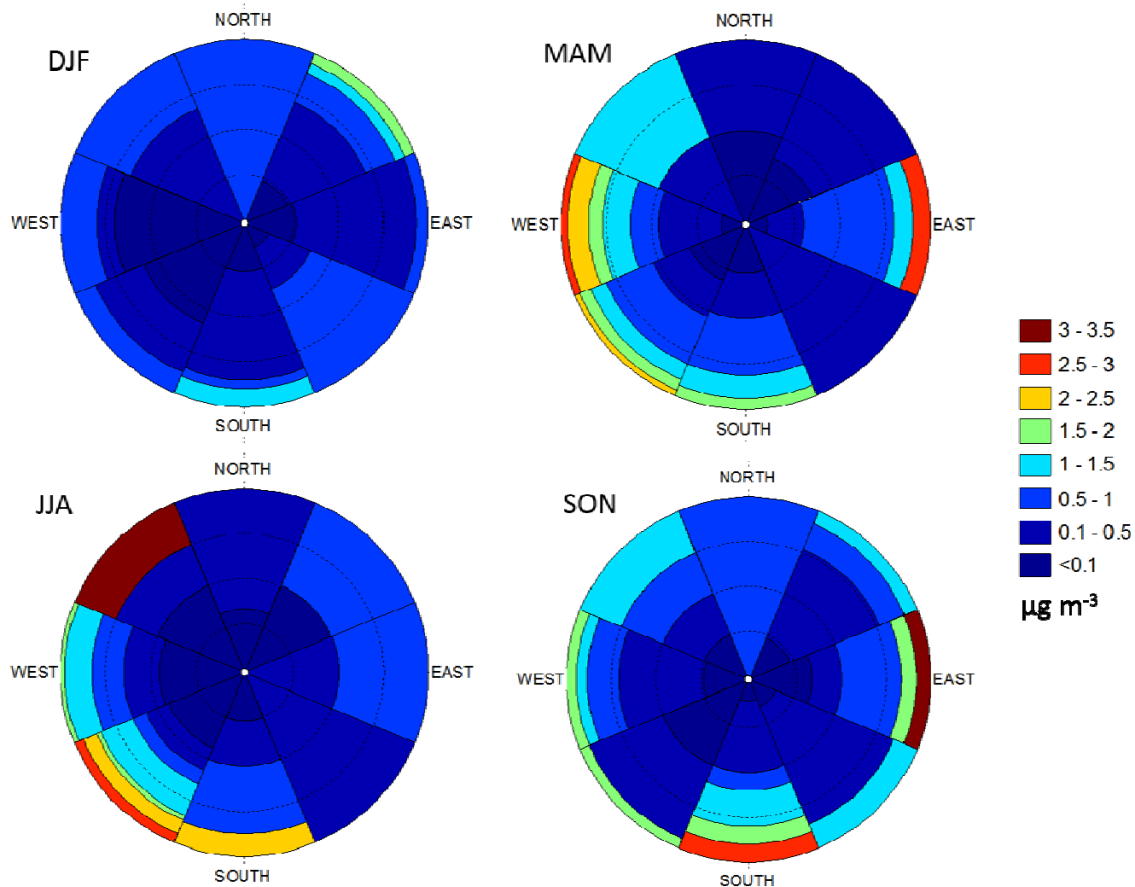
Long term and seasonal trends – aged Sea Salt



Long term and seasonal trends – aged Sea Salt

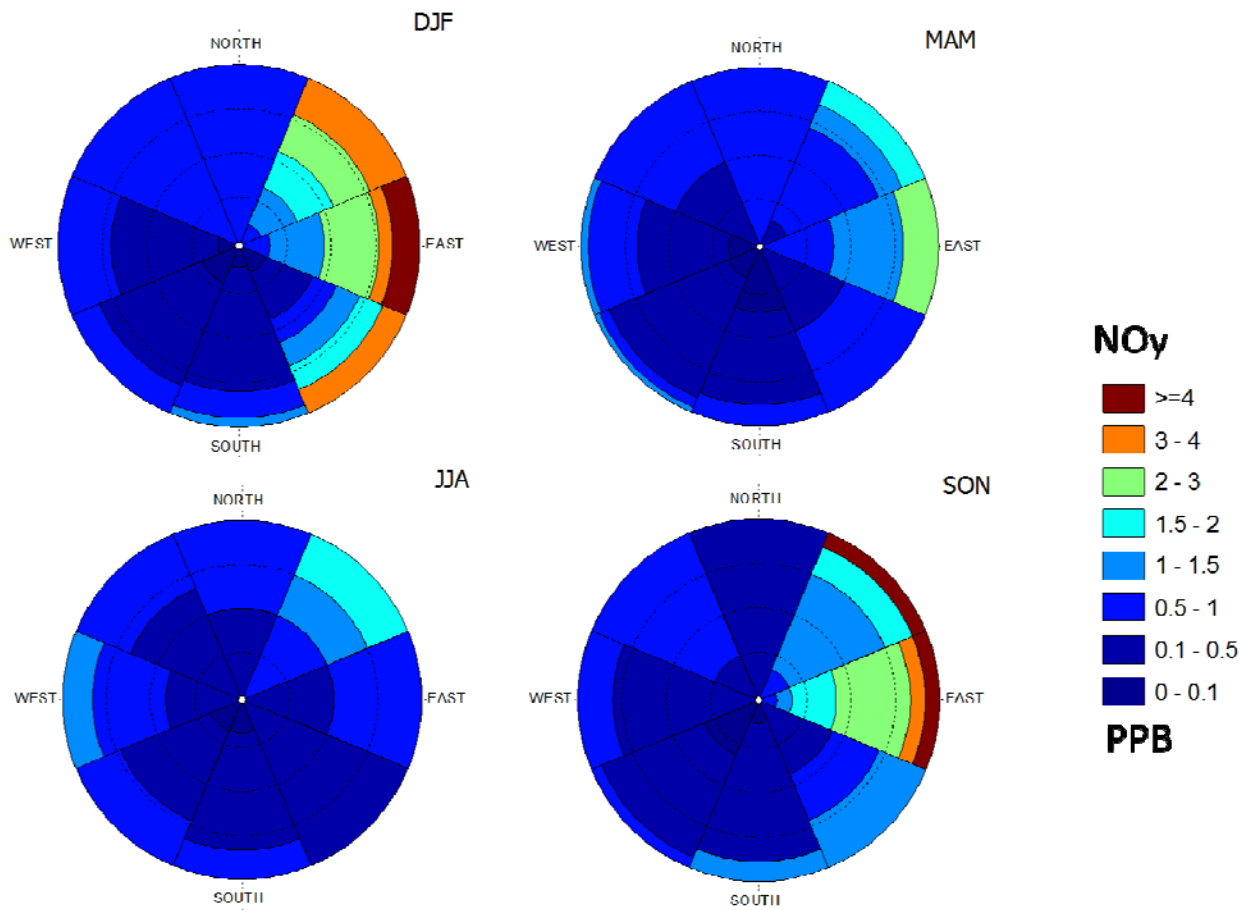


Seasonal wind direction – aged sea salt



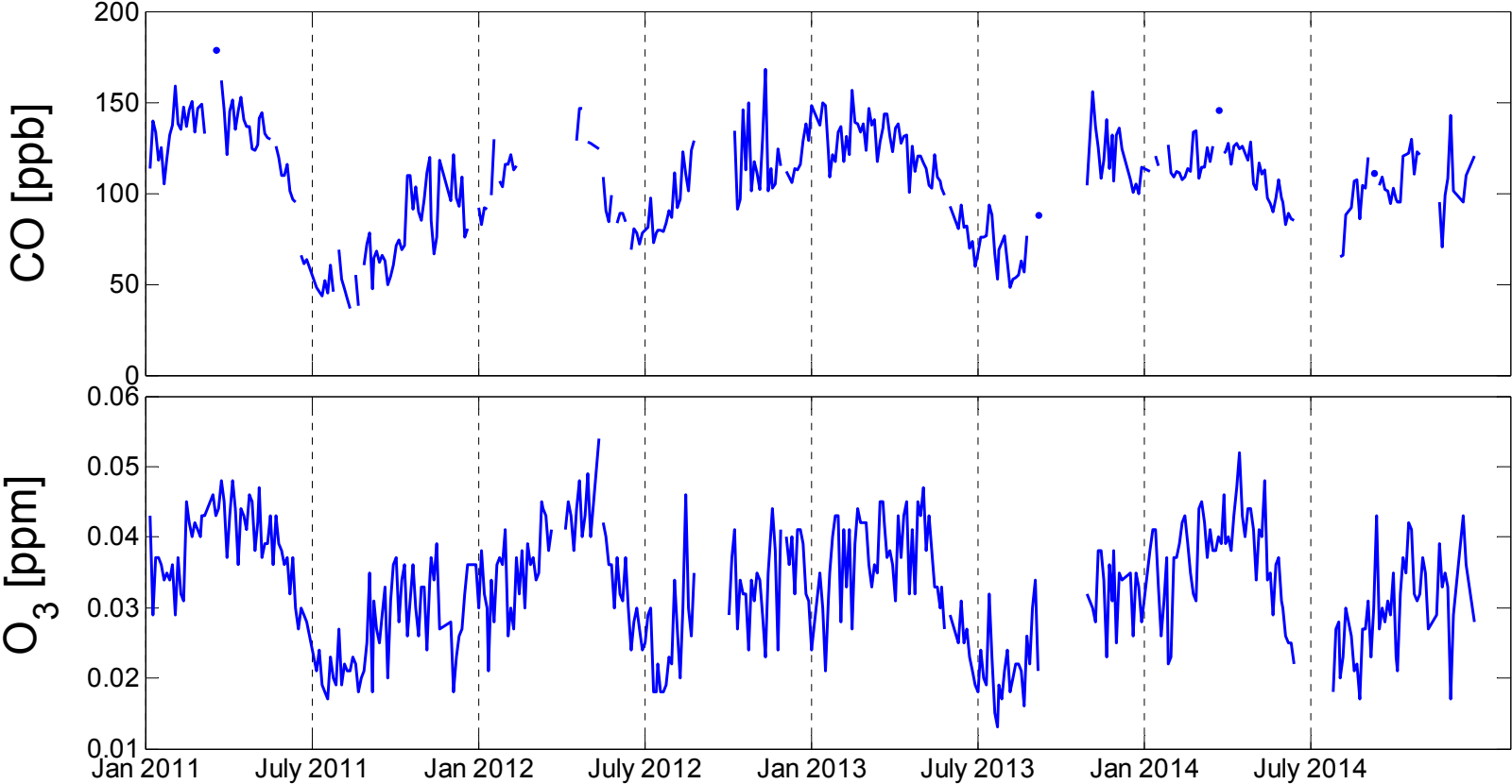
- No clear seasonal correlation
- No clear dependence on wind direction
- The “background” aerosol

Trace Gases – NO/NO_y

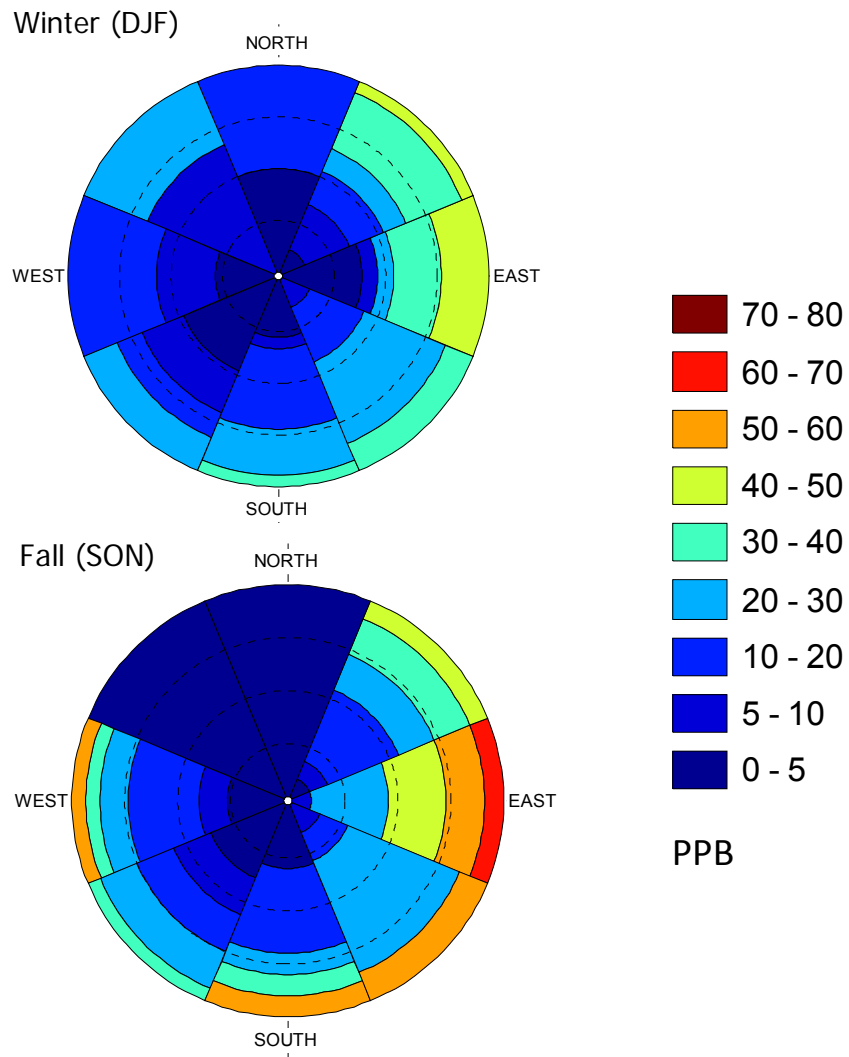


- Similar pattern to BMC source factor
 - Higher concentrations in winter
 - Associated with easterly and northeasterly winds
- Somewhat correlated to BMC ($R^2 = 0.39$)

Trace Gases – O₃ and CO



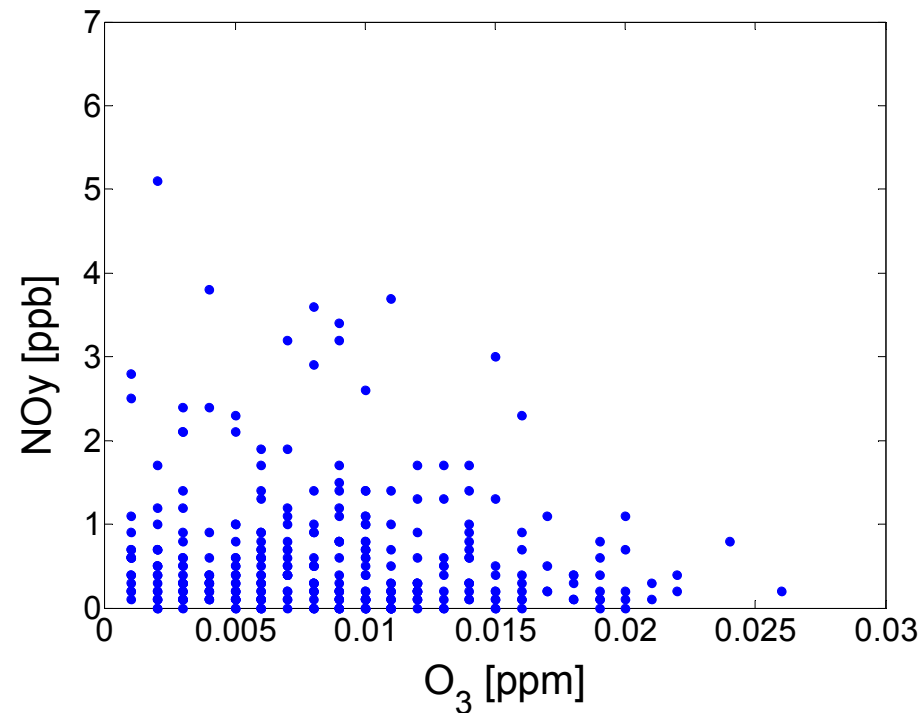
CO residual



- Subtracted minimum monthly value from CO to remove seasonal signal
 - Refer to this as CO residual
- Winter and fall CO residual is enhanced with easterly winds
- No strong correlation with wind direction in summer or spring
- Highest CO residual values observed in the fall
- Most highly correlated with BMC factor ($R^2 = 0.25$)

SO₂ and O₃

- Found no relationship between SO₂ and any source factors
- SO₂ concentrations are almost always less than 1 PPB
- No relationship between O₃ residual and source factors
 - O₃ residual is anti-correlated to NO_y ($R^2 = 0.04$)



Summary

- Marine emissions (RFO) are a dominant source of summer PM2.5
 - Source contribution to PM2.5 has decreased since 2011
 - If low sulfur fuel regulations explain decrease, than chemical fingerprint is likely not the same between 2011 and the following years
- Biomass combustion (BMC) and sea salt are the dominant winter and fall contributors to PM2.5
 - Sea salt is primary pollutant when winds are westerly
 - BMC is dominant when winds are easterly
- Dust is a minor contributor with highest concentrations in spring
 - Associated wind direction is northeasterly (not sure of source)
- Aged aerosol (cloud processed) is a mix of sea salt and ship emissions, indicating older, processed marine aerosol

Acknowledgements

- Environmental Protection Agency (Grant # 00J53301)
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