

Quantification of SO₂ emission rates of the Kīlauea volcano in Hawaii using S5P-TROPOMI satellite measurements

Satellite Remote Sensing Group

Adrian Jost, Steffen Beirle, Steffen Dörner, Christian Borger, Simon Warnach and Thomas Wagner

Max Planck Institute for Chemistry, Mainz, Germany

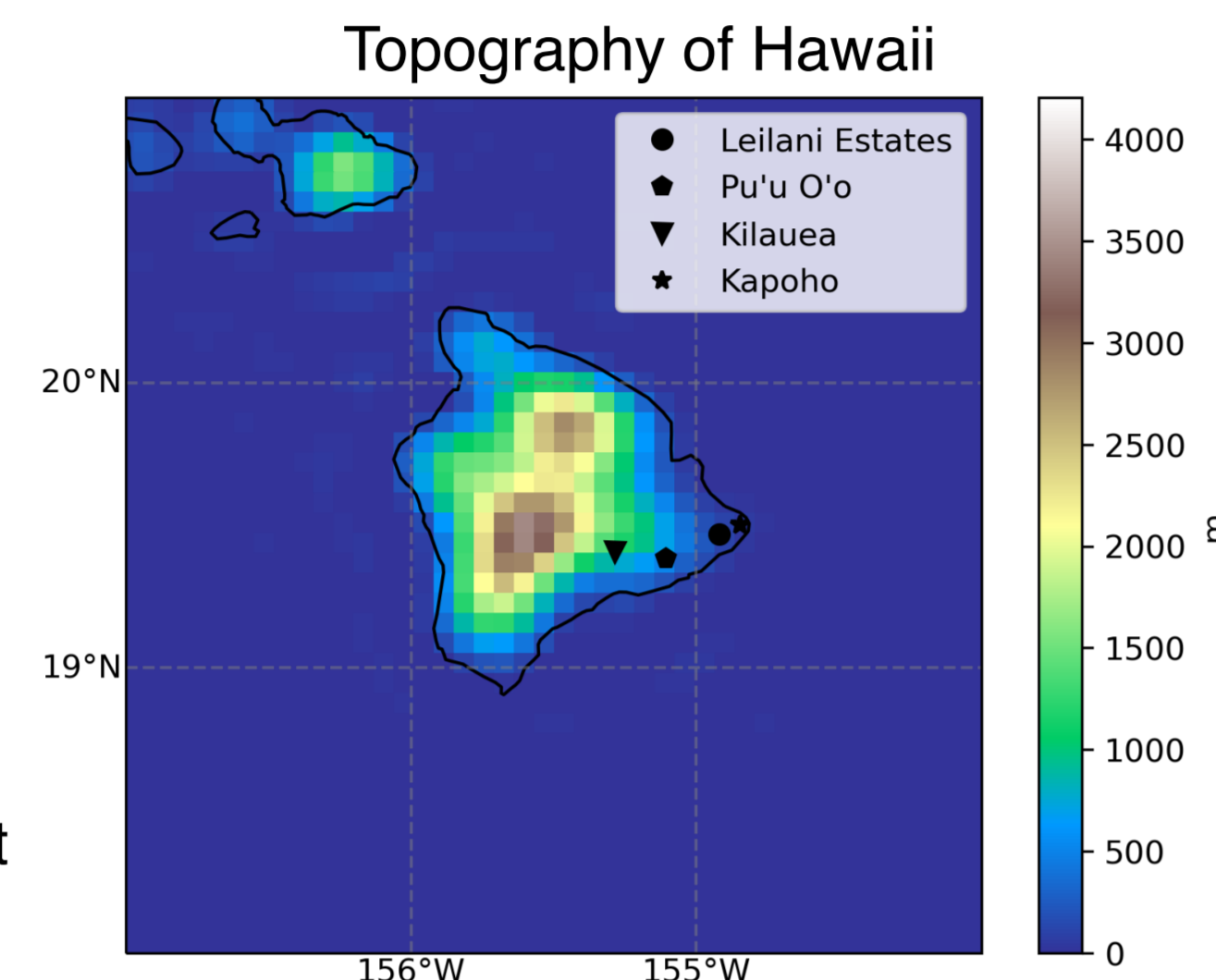
Contact: adrian.jost@mpic.de



Motivation

- SO₂ affects climate and the environment on regional to global scales as well as atmospheric chemistry
- SO₂ is one of the main components of volcanic gases
- TROPOMI provides high spatial resolution measurements of 3.5x7 km
- applying the divergence to the SO₂ flux yields information about the source

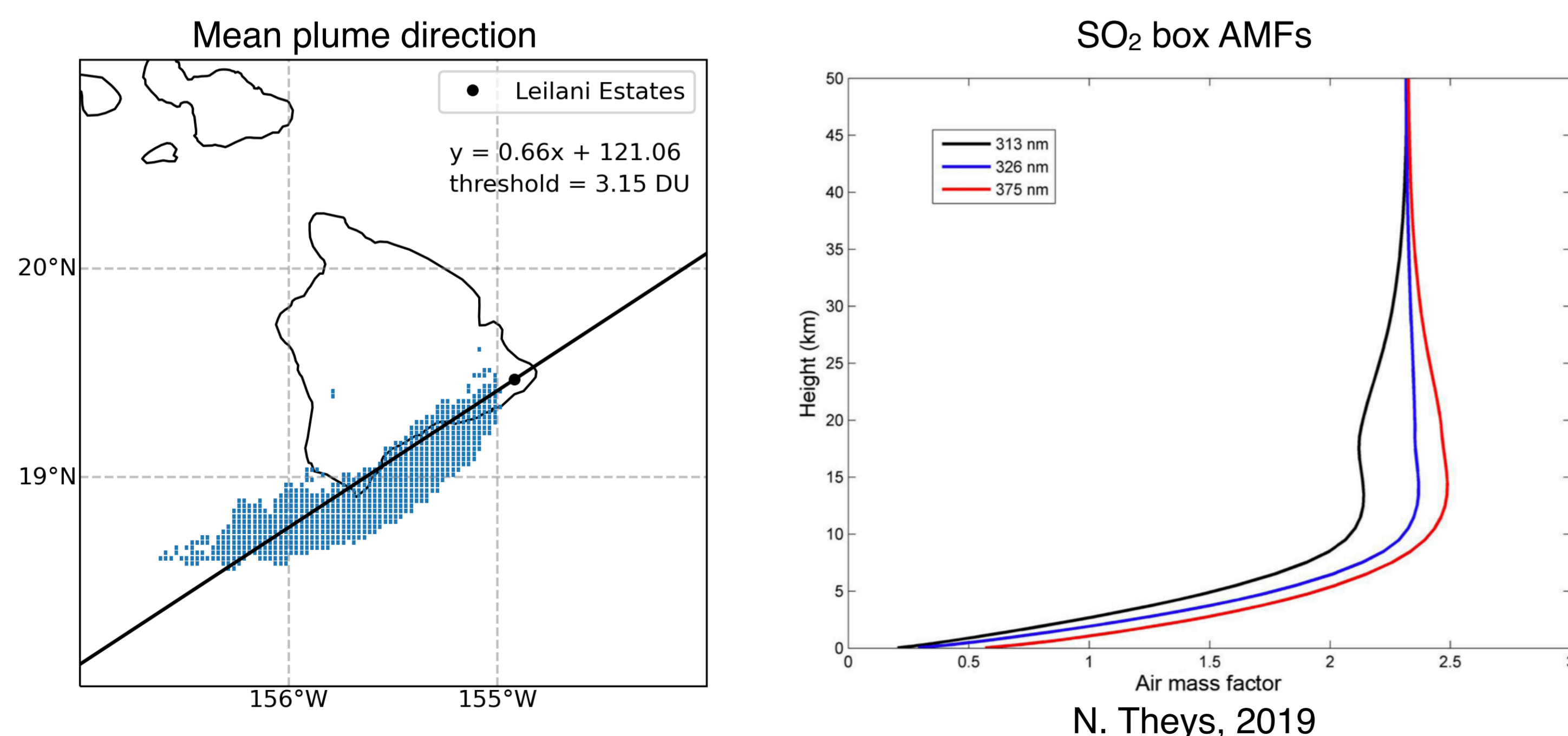
The location and strength of SO₂ emissions from Kilauea are determined for the 2018 Lower East Rift Zone eruption (beginning of May to the end of July) by using the divergence approach.



Algorithm

- selection of two different SO₂ datasets:
 - SCDs of the final selected fitting window from the multiple windows algorithm (NORMAL)
 - SCDs of fitting window 3 only, suitable for high SO₂ columns (WIN3)
- destriping and conversion of slant column densities to vertical column densities using SO₂ box AMFs (N. Theys, 2017)
- filtering of the TROPOMI SO₂ data by omitting high shielding clouds
- gridding of the data to a fine grid with 0.025° horizontal resolution
- interpolation of wind fields to constant heights above sea level
- comparing plume direction with wind direction at different altitudes to find an approximate plume height (plume at approximately 2 km)
- application of the divergence method (S. Beirle, 2019)
- calculation of emission rates for each individual pixel and subsequent summation of pixels around Leilani Estates (source of emissions)

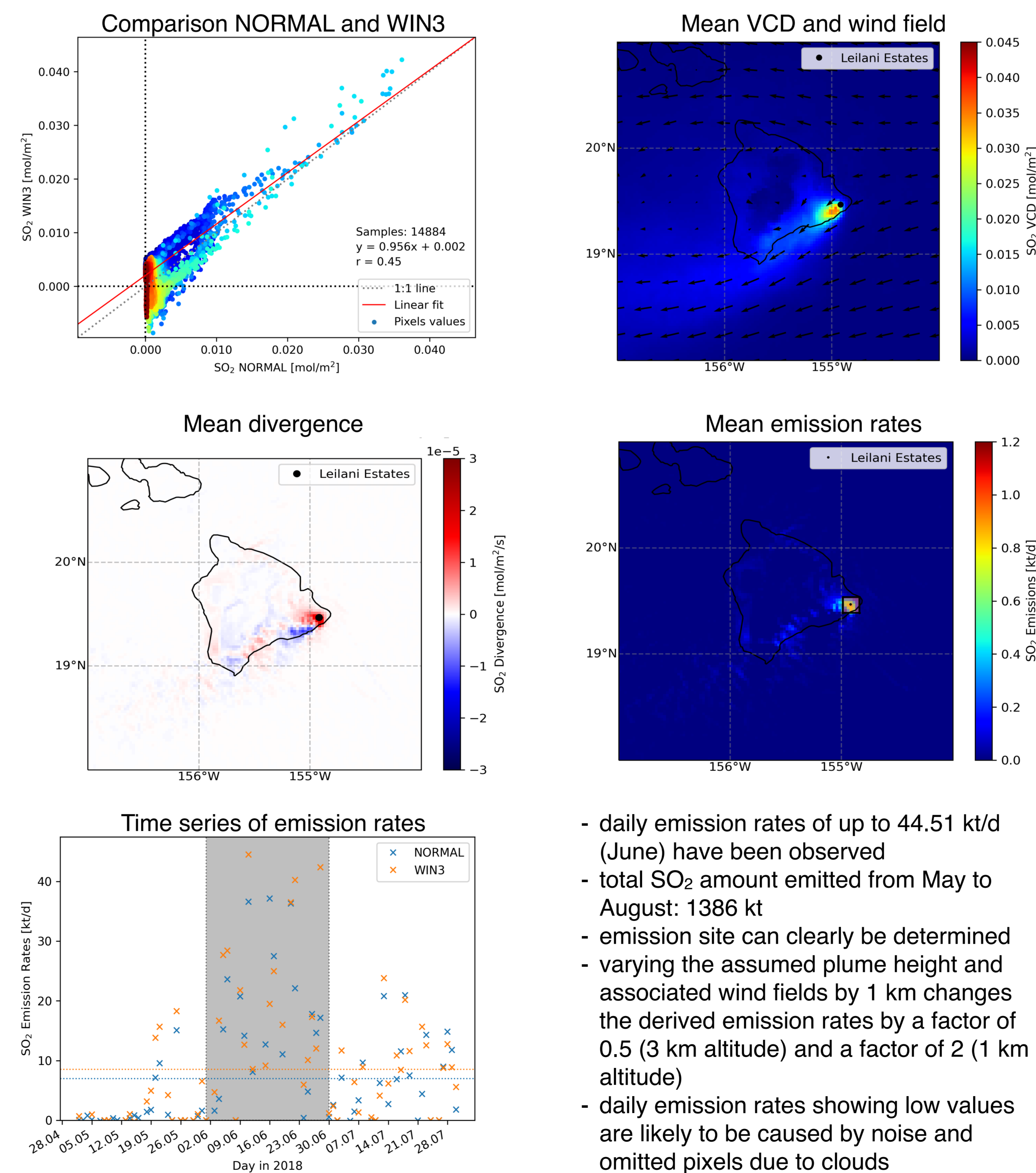
Results have been obtained for single day observations taking only one orbit into account, as well as for the temporal mean SO₂ columns combining three months worth of TROPOMI observations.



Divergence Method

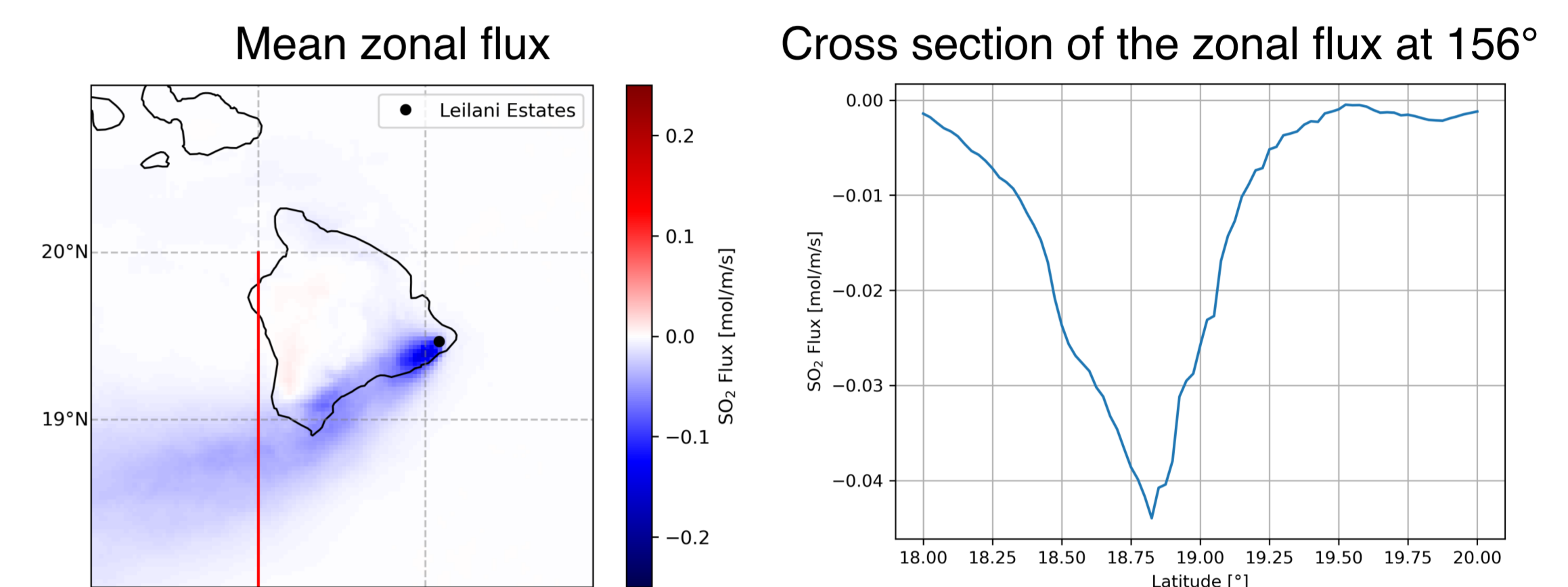
- SO₂ flux is given by:
 $F = \text{VCD} \times w$
- divergence of the horizontal flux yields the sources and sinks of SO₂:
 $D = \nabla(\text{VCD} \times w) = E - S$
- divergence is calculated numerically
- assumption of steady state
- no a priori lifetime is needed

Results

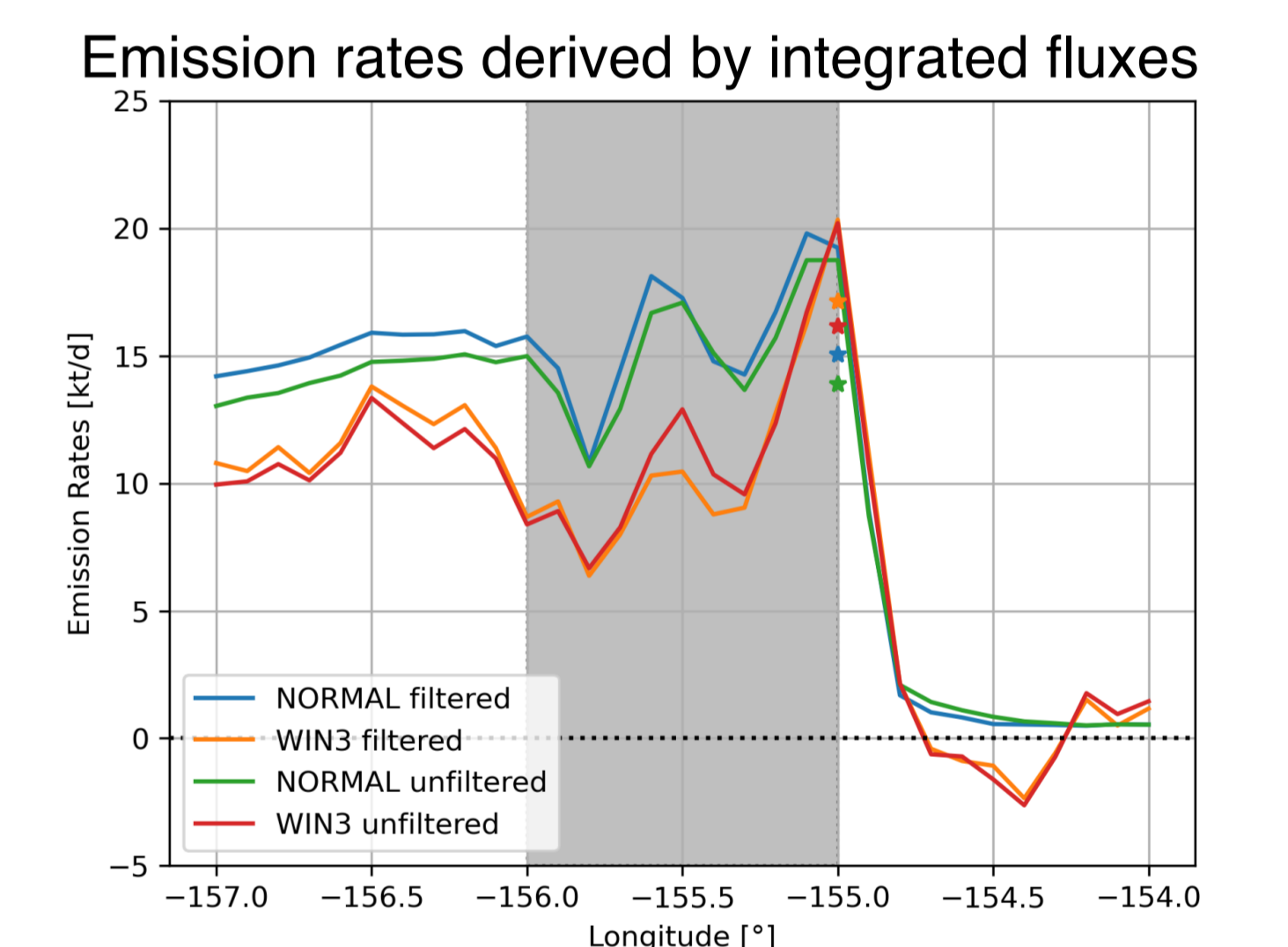


- daily emission rates of up to 44.51 kt/d (June) have been observed
- total SO₂ amount emitted from May to August: 1386 kt
- emission site can clearly be determined
- varying the assumed plume height and associated wind fields by 1 km changes the derived emission rates by a factor of 0.5 (3 km altitude) and a factor of 2 (1 km altitude)
- daily emission rates showing low values are likely to be caused by noise and omitted pixels due to clouds

Alternative Approach



- SO₂ retrieval for dense plumes can be very challenging and uncertain
- diluted plume for cloud-free conditions can be investigated over the Pacific
- calculating the mean flux of a sequence of cross sections and multiplying by the distance from the emission site yields the integrated flux
- the divergence method yields slightly lower values for the mean emission rates



Results obtained by the divergence method can be verified by examining the evolution of the SO₂ flux with increasing distance.

Discussion

Location and strength of SO₂ emissions can be determined for a high spatial resolution and the results are verified by the flux integration method. Obtained emission rates are much lower than for ground-based observations due to a different AMF (C. Kern, 2020).

- plume height holds a very large uncertainty and therefore has a decisive impact on the wind fields and the AMF
- poor statistics for individual orbits makes multi-day averaging necessary
- further investigation of the variables leading to high uncertainties
- application on a global scale