

Derivation of vertical profiles of SO₂ in the Venus cloud layer by the Akatsuki radio occultation measurements.

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Introduction: On Venus, sulfuric acid (H₂SO₄) clouds exist at the altitudes from 45 to 70 km, covering the entire planet. This thick cloud layer plays a key role in controlling the heat budget in Venus atmosphere. One of the missing information in the cloud layer is the vertical distribution of the sulfur dioxide (SO₂), which is one of the main chemical compounds in forming the H₂SO₄ clouds.

The previous observations of the SO₂ vertical distribution at the cloud layer were limited to the in-situ measurements by the VEGA probes [1]. The two probes measured the vertical profiles of the SO₂ mixing ratio from the surface to the altitude of 60 km. In addition to the in-situ measurements, radio occultation (RO) measurements of the Venus Express derived the mean SO₂ mixing ratio at 51-54 km [2].

Extending the method adopted in the Venus Express RO measurements, we have derived the vertical profiles of SO₂ mixing ratio in the cloud layer (50-55 km) by the RO measurements in the Japanese Venus orbiter, "Akatsuki".

Data and method: We analyzed the Akatsuki RO data obtained at the low latitudes (40°S-40°N) during 5 Earth years (2016-2021). In the RO measurements, we utilize the absorption of the radio waves transmitted from the spacecraft to the Earth, which pass through the Venus atmosphere. Time series data of received intensity can be obtained, and the distribution of mixing ratio of trace substances can be obtained from the time variation of radio wave intensity with the attenuation of the radio waves.

In the frequency domain used in the Akatsuki RO measurements (8.4 GHz), the gaseous species including CO₂, N₂, H₂SO₄ vapor and SO₂ contribute to the attenuation of the radio waves in the Venus atmosphere. In addition to the absorption by the constituents in the Venus atmosphere, the defocusing loss, the directional error of the transmitter antenna and Earth's atmosphere also contribute to the attenuation. We removed those effects from the original radio wave's intensity to obtain the residual attenuation. The concentrations of H₂SO₄ vapor and SO₂ are then derived by subtracting the attenuations caused by CO₂ and N₂. Finally, we assume that H₂SO₄ vapor is not supersaturated in the Venus cloud layer and any attenuations which exceed the saturation curve of H₂SO₄ vapor attribute to the absorption by SO₂.

Error analysis: We conducted the error analysis of the SO₂ mixing ratio retrieved as follows:

Effect of the interplanetary plasma and Earth's atmosphere. As mentioned above, the radio waves

emitted from the spacecraft orbiting Venus is also attenuated by the Earth's atmosphere. The significance of this effect was estimated by analyzing the short time-scale variations appearing in the time series of the received radio wave intensity before/after the occultation. We found that the error of the SO₂ mixing ratio possibly exceeds 100% at 50-55 km due to those effects, suggesting that the precision of the individual observation was not high. Therefore, we averaged all the vertical profiles of SO₂ to improve the signal-to-noise ratios.

Uncertainty of the H₂SO₄ saturation mixing ratio. The H₂SO₄ saturation mixing ratio depends not only on temperature but also on the mass concentration and the size of the cloud particles. The higher the mass concentration and the smaller the particle size, the greater the H₂SO₄ saturation mixing ratio. Numerical models suggest that the mass concentration of H₂SO₄ is higher than 98% at 50 km and nearly 90% at 55 km [3]. The radius of the Venus cloud particles ranges from submicron (mode 1) to several microns (mode 3) [4]. We estimated the uncertainty of the H₂SO₄ saturation mixing ratio were 25-50%.

Uncertainty of the absorptivity of CO₂, N₂ and H₂SO₄ vapor. We examined the uncertainties in the absorptivity of CO₂ and N₂ combined and H₂SO₄ vapor. These were ~3 % and ~10 %, respectively. Considering all the uncertainties in the *absorptivity of the gases*, SO₂ mixing ratios derived in this study have an uncertainty of ~30 %.

Result of the SO₂ mixing ratio retrieved: The averaged vertical profile showed the SO₂ mixing ratio of about 300 ppmv and 50 ppmv at the altitude of 50 km and 55 km, respectively, decreasing with increasing altitude. Those are consistent with the previous observations of the VEGA probes and the Venus Express RO measurements. Additionally, we found a local time dependence of the SO₂ mixing ratio in the cloud layer, which was smaller in the daytime than in the nighttime, being similar with the previous observations conducted above the cloud layer [5]. The long-term variations were not detected in the present study.

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References: [1] Bertaux et al. (1996) J. Geophys Res 101(E5), 12709–12745. [2] Oschlisniok et al.

(2021) *Icarus*, Volume 362, 2021. [3] Hashimoto and Abe (2001) *J. Geophys. Res.*, 106(E7), 14675– 14690. [4] Knollenberg and Hunten, *J. Geophys. Res.* 85, 8039 (1980) [5] Marcq et al. (2020) *Icarus*, Volume 335, 2020.