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On recent (2008–2012) stratospheric aerosols observed by lidar over Japan

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Abstract

An increase in stratospheric aerosols caused by the volcanic eruption of Mt. Nabro $(13.37^{\circ} \text{ N}, 41.70^{\circ} \text{ E})$ on 12 June 2011 was first detected by lidar at Tsukuba $(36.05^{\circ} \text{ N}, 140.13^{\circ} \text{ E})$ and Saga $(33.24^{\circ} \text{ N}, 130.29^{\circ} \text{ E})$ in Japan. The maximum backscattering ratios at a wavelength of 532 nm were 2.0 at 17.0 km on 10 July 2011 at Tsukuba and 3.6 at 18.2 km on 23 June 2011 at Saga. The maximum integrated backscattering coefficients (IBCs) above the first tropopause height were $4.18 \times 10^{-4} \text{ sr}^{-1}$ on 11 February 2012 at Tsukuba and $4.19 \times 10^{-4} \text{ sr}^{-1}$ on 23 June 2011 at Saga, respectively.

A time series of lidar observational results at Tsukuba have also been reported from January 2008 through May 2012. Increases in stratospheric aerosols were observed after the volcanic eruptions of Mt. Kasatochi (52.18° N, 175.51° E) in August 2008 and Mt. Sarychev Peak (48.09° N, 153.20° E) in June 2009. The yearly averaged IBCs at Tsukuba were 2.60 × 10⁻⁴ sr⁻¹, 2.52 × 10⁻⁴ sr⁻¹, 2.45 × 10⁻⁴ sr⁻¹, and 2.20 × 10⁻⁴ sr⁻¹ for 2008, 2009, 2010, and 2011, respectively. These values were about twice the IBC background level (1.21 × 10⁻⁴ sr⁻¹) from 1997 to 2001 at Tsukuba. We briefly discuss the influence of the increased aerosols on climate and the implications for analysis of satellite data.

1 Introduction

Stratospheric aerosols play important roles in climate regulation and atmospheric
 chemistry. The effects of the aerosols produced by the Pinatubo eruption is a good example. The volcanic eruption of Mt. Pinatubo (15.14° N, 120.35° E) on 15 June 1991 injected huge amounts of SO₂ and ash into the stratosphere. The Volcanic Explosivity Index (VEI) was 6 (Smithsonian Institution, 2012). The eruption injected into the stratosphere an amount of SO₂ estimated to be about 20 Tg, almost three times the 1982 El Chichón eruption (Bluth et al., 1992). The injected SO₂ was oxidized to sulfuric acid particles through homogeneous nucleation (Wu et al., 1994).





Read et al. (1993) estimated the e-folding decay time of SO₂ to be 33 days. The first increase of aerosols from the Pinatubo eruption was observed at an altitude of 15.7 km over Tsukuba (36.05° N, 140.13° E) on 28 June 1991. The Pinatubo aerosol particles were effectively transported from tropical regions into northern mid-latitudes during

- ⁵ fall through spring with planetary wave activity. The maximum backscattering ratio observed at a wavelength of 532 nm was 14.1 at 22.7 km over Tsukuba on 29 November 1991. The maximum value of the integrated backscattering coefficient (IBC) above the first tropopause height was $7.1 \times 10^{-3} \text{ sr}^{-1}$ over Tsukuba on 22 February 1992 (Uchino et al., 1995).
- ¹⁰ The stratospheric aerosol surface area increased after the Pinatubo eruption (Jäger et al., 1995; Uchino, 1996), and severe ozone loss occurred in 1992 and 1993 because of heterogeneous chemical reactions on aerosol surfaces in the presence of high concentrations of anthropogenic chlorine and bromine (Hofmann et al., 1994; Kondo et al., 1995; WMO, 1995; Solomon et al., 1996).
- The maximum net (thermal minus solar) radiative forcing from the 1991 Pinatubo eruption was about -3 W m⁻² (Hansen et al., 2005). Global lower stratospheric (30–100 hPa) temperature anomalies increased after the eruption, and global tropospheric (300–850 hPa) temperature anomalies decreased after the eruption in spite of the warm ENSO episode in 1991/1992 (Kawamata et al., 1992). Global tropospheric tem-
- ²⁰ peratures generally increase after a warm ENSO episode. For two years following major volcanic eruptions, global mean surface temperatures decrease by 0.1–0.2 °C, and by 0.3 °C during the summer in the latitude band 30–60° N (Robock and Mao, 1995). A model simulation of the effects of the 1991 Pinatubo eruption predicted a decrease in the global surface temperature by about 0.5 °C in September, October, and November 1992. in agreement with observations during that time (Hansen et al., 1996).

In contrast, warm surface temperatures were recorded over Europe, Siberia, and North America, while cooling occurred over western Asia in the winters after the three major volcanic eruptions of Mt. Agung in 1963, Mt. El Chichón in 1982, and Mt. Pinatubo in 1991. Volcanic aerosols produce a stronger westerly jet in the stratosphere





during the winter because they increase the temperature gradient between the equator and high latitudes. The stronger stratospheric polar night jet extends into the troposphere through interactions with planetary waves, and changes in tropospheric circulation induce a stronger polar vortex and equatorward propagation of waves (Kodera,

⁵ 1994). As a result, warm tropospheric temperature anomalies occur in the winter in the Northern Hemisphere after major volcanic eruptions in the tropics.

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The IBC of the Pinatubo aerosols decayed with e-folding times of 1.14, 1.29, and 1.37 yr over Tsukuba and Naha (26.2° N, 127.7° E) in Japan and over Lauder (45.0° S, 169.7° E) in New Zealand, respectively. The IBC over Tsukuba varied in a clearly seasonal manner, with a maximum in winter and early spring and a minimum in summer.

The IBC over Tsukuba reached the background level in October 1997 (Nagai et al., 2010).

Since about 2000, an increase of 4–7% per year in the IBC has been detected within the 20–30 km altitude range at both Mauna Loa, Hawaii (19°N), and Boulder,

- ¹⁵ Colorado (40° N) (Hofmann et al., 2009). Likewise, after the IBC over Lauder reached a minimum between 1997 and 2000, it increased 3.8% per year from 2000 to 2009 (Nagai et al., 2010). Based on some satellite data, the stratospheric aerosol optical thickness (AOT) increased after 2000 as the result of a series of moderate but increasingly intense volcanic eruptions (Vernier et al., 2011). In fact, increases in strato-
- ²⁰ spheric aerosols were reported from lidar observations after the volcanic eruptions of Mt. Kasatochi (52.18° N, 175.51° E) in August 2008 (Bitar et al., 2010) and Mt. Sarychev Peak (48.09° N, 153.20° E) in June 2009 (Uchino et al., 2010; O'Neill et al., 2012).

In this paper we report the first observational results of stratospheric aerosols in the year following the volcanic eruption of Mt. Nabro (13.37° N, 41.70° E) in June 2011

at two lidar sites in Tsukuba and Saga (33.24° N, 130.29° E), Japan. These two lidar sites are prioritized validation sites for studying the influence of aerosols and thin cirrus clouds on column-averaged dry air mole fractions of carbon dioxide (XCO₂) and methane (XCH₄) derived from data collected by the Greenhouse gases Observing SATellite (GOSAT) (Yoshida et al., 2011; Morino et al., 2011; Uchino et al., 2012).





GOSAT was launched on 23 January 2009. At Saga, lidar observations started in March 2010. Next, we present lidar observational results from January 2008 to May 2012 over Tsukuba. Finally we discuss briefly the influence of the recent increase in stratospheric aerosols on GOSAT products and compare their impact on climate to the 1991 Pinatubo eruption.

2 Lidar instruments and data analysis

The compact lidars installed at Tsukuba and Saga were two-wavelength polarization lidar systems (Table 1), the fundamental and second harmonic having wavelengths of 1064 nm (λ_1) and 532 nm (λ_2), respectively. Backscattered photons from the atmosphere were collected by one or two Schmidt Cassegrain type telescopes. A polarizer divided photons at λ_2 into components parallel (P) and perpendicular (S) to the transmitted laser polarization plane. The received photons were converted to electrical signals by an avalanche photodiode (APD, C30956EH) at λ_1 . At λ_2 , three or five photomultiplier tubes (PMTs, R3234-01) were used to simultaneously obtain high-dynamicrange signals from near the surface to an altitude of ~ 40 km. Transient recorders used a 12-bit analog-to-digital (A/D) converter and a photon counter (TR 20-160) to process the output signals of the APD and PMTs. Because the APD signals were noisy above altitudes of about 20–25 km, we used only lidar data at λ_2 for stratospheric aerosols.

The backscattering ratio R is defined as

 $_{20} \quad R = (BR + BA)/BR,$

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where *BR* and *BA* are the molecular and aerosol backscattering coefficients, respectively. We derived backscattering ratio profiles with an inversion method (Fernald, 1984). The lidar ratio *S* (particle extinction to backscatter ratio) is dependent on the stratospheric aerosol size distribution and refractive index, and equalled 20–60 sr at 532 nm during 1979–1999 (Jäger and Deshler, 2002, 2003). The lidar ratio was small just after the major volcanic eruptions of El Chichon in 1982 and Pinatubo in 1991, but



(1)



it equals about 50 sr for usual stratospheric aerosols. We assumed that the lidar ratio equalled 50 sr for the moderate volcanic eruptions of Kasatochi in 2008, Sarychev in June 2009 and Nabro in June 2011. We used the nearest operational radiosonde data to calculate the atmospheric molecular density. The radiosonde sounding stations are

- Tateno (36.05° N, 140.13° E) and Fukuoka (33.58° N, 130.38° E) for Tsukuba and Saga, respectively. We used the 1976 US Standard Atmosphere model above balloon burst altitudes (US Committee on Extension of the Standard Atmosphere, 1976). The lidar backscattered signal was interactively normalized to unity around 25–33 km, where aerosol-free conditions could be assumed.
- ¹⁰ We obtained IBCs by summing up *BA*s from the first tropopause height to an altitude of 33 km. When cirrus clouds appeared above the tropopause, we set the lower limit of the integration to just above the altitude of the cirrus clouds. If the signal-to-noise ratio at higher altitudes was not good enough for integration purposes, the upper limit of the integration was decreased to a lower altitude where the signal-to-noise ratio was acceptable (Nagai et al., 2010)

The total linear depolarization ratio (δ) is defined as

 $\delta = S/(P+S) \cdot 100\,(\%)$

where *P* and *S* are the parallel and perpendicular components of the backscattered signals. The particle depolarization δ_{p} is obtained from the equation

 $\delta_{\rm p} = (\delta \cdot R - \delta_{\rm m})/(R - 1) \cdot 100(\%)$

(3)

where δ_m is the depolarization ratio of atmospheric molecules (Sakai et al., 2003). We adopted a vertical resolution of 150 m in the following analysis.

25 **3** Observational results over Tsukuba and Saga after the 2011 Nabro eruption

The Nabro volcano erupted in Eritrea on 12 June 2011. The volcanic ash was detected at 10:45 UTC on 13 June by the Moderate Resolution Imaging Spectrometer



(2)



(MODIS) on the Aqua satellite (NASA, 2012). The first SO₂ associated with the eruption was measured on 12 June by the Infrared Atmospheric Sounding Interferometer (IASI), and continued emissions were observed for weeks. The total mass of SO₂ measured by IASI was on the order of 1.5 Tg (Clarisse et al., 2012). Over Tsukuba, new aerosol layers with double peaks were observed on 20 June 2011 about 8 days after the eruption (Fig. 1). The peak values of *R* were 1.58 and 1.32 at 16.0 and 16.4 km, respectively. The values of δ and δ_p were 1.25% and 3.4%, respectively, at 16.0 km and 1.94% and 7.9%, respectively, at 16.4 km. Non-spherical ash particles were probably included in the layers with sulfuric acid particles that were produced from SO₂ through chemical reactions. Non-spherical particles were also present in the lower region of the aerosol layer on 12 September ($\delta_p = 4.7\%$ at 17.0 km). The maximum backscattering ratio (R_{max}) of 2.0 was observed at 17.0 km on 10 July 2011.

Over Saga, new stratospheric aerosols with double peaks were detected on 23 June 2011. Peak values of *R* were 2.27 and 3.68 at 17.2 and 18.2 km, respectively. The values of δ_p were 0.2% and 0.8% at 17.2 and 18.2 km, respectively. In this case, aerosols were probably composed of spherical particles because δ_p was very small. However, some non-spherical particles were also seen in the lower regions of the layers on 29 August ($\delta_p = 3.6\%$ at 16.6 km) and 24 September ($\delta_p = 4.0\%$ at 16.6 km). In the 1991 Pinatubo eruption, non-spherical particles were present in the lower stratosphere

²⁰ for at least six months (Nagai et al., 1993) because the Pinatubo ash particles were injected into higher altitudes than the Nabro ash particles.

We used the National Centers for Environmental Prediction (NCEP)/National Center for Atmospheric Research (NCAR) reanalysis data (Kalnay et al., 1996) and the Meteorological Data Explorer (METEX), developed by Jiye Zeng at the Centre for Global

Environmental Research (CGER) in the National Institute for Environmental Studies (NIES) to calculate isentropic forward trajectories of 36 air parcels that originated from a square of ±1° surrounding the Nabro volcano at altitudes of 16, 17, and 18 km. The calculation simulated the trajectories of the air parcels for ten days beginning at 23:00 UTC on 12 June 2011. Only some of the parcels that originated at 17 km (potential tempera-



ture of 384.3 K) over Mt. Nabro were transported to ~ 16 km over Tsukuba on 20 June, a result that was consistent with lidar observations as shown in Fig. 1. The air parcels moved eastward around the northern part of the Tibetan high-pressure ridge (Fig. 3). The composite image of maximum observed SO₂ columns in Fig. 12 of Clarisse et al.

- (2012) also shows this feature. We confirmed that the backward trajectory of an air parcel from Tsukuba (16 km, 13:00 UT on 20 June 2011) arrived at a point (16.7 km, 14.62° N, 33.42° E) near Mt. Nabro on 23:00 UTC on 12 June. Therefore, new aerosol layers observed over Japan in late June 2011 could have originated from the Nabro eruption on 12 June.
- Figure 4 shows the time variation of IBC (pink solid diamond) and first tropopause height (blue open circle) over Tsukuba (upper panel) and Saga (lower panel) from June 2011 to May 2012. Over Tsukuba, the largest values of the IBC were $\sim 3.0 \times 10^{-4} \, \text{sr}^{-1}$ in summer and $\sim 4.0 \times 10^{-4} \, \text{sr}^{-1}$ in winter. The maximum IBC was $4.18 \times 10^{-4} \, \text{sr}^{-1}$ on 11 February 2012. In general the IBC increased when the tropopause height decreased. Over Saga, the maximum IBC was $4.19 \times 10^{-4} \, \text{sr}^{-1}$ on 23 June 2011, the day of the first arrival of the Nabro aerosols. Then the IBC decreased quickly within a week, but increased again in late July. The IBC then decreased gradually from August
- to December 2011, except for a brief peak larger than $\sim 3.5 \times 10^{-4} \text{ sr}^{-1}$ on 24 and 25 November. The IBC increased again in January and February 2012. The mean value of the IBC over Saga was $1.86 \times 10^{-4} \text{ sr}^{-1}$ from June 2011 to May 2012.

4 Time variation of stratospheric aerosols over Tsukuba from January 2008 to May 2012 and discussion

Mt. Kasatochi in the Aleutian Islands erupted on 7 and 8 August 2008, and the VEI was 4 (Smithsonian Institution, 2012). The Ozone Monitoring Instrument (OMI) on NASA's Aura satellite tracked a dense cloud that contained about 1.5 Tg of SO₂. The SO₂ clouds spread over the Arctic and eastward across the United States and Canada (NASA, 2012). Over Halifax (44.64° N, 63.59° W) in Canada, aerosols from the vol-



canic plume were detected with lidar one week after the eruption and for the next four months thereafter (Bitar et al., 2010). Over Tsukuba, stratospheric aerosols produced from those SO_2 gases were detected at 17.3 km and 16.0 km on 2 September and at 18.7 km and 17.3 km on 16 September, about one month after the eruption (Fig. 5). Clear peaks of *R* were also seen on 4 and 21 October, but subsequently those peaks were ambiguous. Obvious atratespheric aerosols from the Kasatashi aruption

peaks were ambiguous. Obvious stratospheric aerosols from the Kasatochi eruption were also observed from 10 September to 13 October over Ryori (39.03° N, 141.82° E) (Sakashita et al., 2009).

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Mt. Sarychev Peak erupted on 12 June 2009, and the VEI was 4 (Smithsonian Institution, 2012). A new aerosol layer was observed at 20.6 km on 25 June over Tsukuba (Fig. 6). The peak value of *R* was 3.5. Because δ_p was 7%, some non-spherical ash particles were probably included in the layer. Backward trajectory analysis revealed that aerosols in the layer were transported to Tsukuba by easterly winds. Aerosols observed around 14–15 km on 5 July were transported by westerly winds. Enhanced aerosol lay-

ers were also observed over other three lidar sites in Japan (Uchino et al., 2010). Mt. Merapi (7.54° S, 110.44° E), one of Indonesia's most active volcanoes, erupted on October 2010, and the VEI was 4 (Smithsonian, 2012). Shortly thereafter we did not observe enhanced stratospheric aerosols that originated from the Merapi eruption, because noticeable peaks of *R* were not detected.

²⁰ The temporal variation of the IBC over Tsukuba from January 2008 through May 2012 is shown in Fig. 7, with the exception of about two months in 2011 after the Tohoku earthquake off the Pacific Coast of Japan, when lidar data were not obtained. The earthquake occurred in the northern part of Japan on 11 March 2011. After the decay of the Pinatubo aerosols, stratospheric aerosols were at background levels from October

²⁵ 1997 to September 2001 at Tsukuba (Nagai et al., 2010). The annual mean of the IBC for the background aerosols was $1.21 \times 10^{-4} \, \text{sr}^{-1}$. Based on the fit of a sinusoidal function to the data, the amplitude of the seasonal variation was $6.84 \times 10^{-5} \, \text{sr}^{-1}$, with a maximum in February and minimum in August (Fig. 7). According to Deshler et al. (2006), no long-term change in the background concentration of stratospheric aerosols





has occurred over the period 1972–2004, and therefore the background level of the IBC observed over Tsukuba from October 1997 to September 2001 might be similar to the background levels during the period 1972–2004.

Most IBCs from January 2008 through May 2012 in Fig. 7 were larger than those associated with background aerosols during October 1997 through September 2001. The IBCs increased after the volcanic eruptions of Mt. Kasatochi in August 2008 and Mt. Sarychev Peak in June 2009. The total masses of SO₂ from the Kasatochi, Sarychev Peak, and Nabro eruptions were estimated to be 1.6 Tg, 0.9 Tg, and 1.5 Tg, respectively (Clarisse et al., 2012). However, the production rate of stratospheric aerosols depends on the amounts of SO₂ that are injected into the stratosphere. Before the Kasatochi

- on the amounts of SO_2 that are injected into the stratosphere. Before the Kasatochi eruption, the IBC was larger than the background level, an observation consistent with that of Vernier et al. (2011) and possibly due to some other volcanic eruptions in the tropics, including Tavurvur (4.27° S, 152.2° E) on 7 October 2006 and Soufrière Hills (16.72° N, 62.18° W) on 20 May 2006.
- ¹⁵ The yearly averaged IBCs over Tsukuba were $2.60 \times 10^{-4} \text{ sr}^{-1}$, $2.52 \times 10^{-4} \text{ sr}^{-1}$, $2.45 \times 10^{-4} \text{ sr}^{-1}$, and $2.20 \times 10^{-4} \text{ sr}^{-1}$ for 2008, 2009, 2010, and 2011, respectively. Therefore the elevations of the IBCs above background level were $1.39 \times 10^{-4} \text{ sr}^{-1}$, $1.31 \times 10^{-4} \text{ sr}^{-1}$, $1.24 \times 10^{-4} \text{ sr}^{-1}$, and $0.99 \times 10^{-4} \text{ sr}^{-1}$, respectively. The corresponding elevations of the AOTs above background levels were 0.0070, 0.0066, 0.0062, and
- ²⁰ 0.0050, respectively, for an assumed lidar ratio of 50 sr. The corresponding increases of negative radiative forcing (cooling) were roughly 0.18 Wm⁻², 0.17 Wm⁻², 0.16 Wm⁻² and 0.13 Wm⁻², respectively, based on a conversion factor of 25 Wm⁻² from AOT to radiative forcing (Hansen et al., 2005; Solomon et al., 2011). These values are not small compared to the positive radiative forcing (heating) caused by increases in at-
- ²⁵ mospheric CO₂, which has averaged about 0.28 Wm⁻² over the decade since 2000 (Solomon et al., 2011; NOAA, 2012). The average AOT for the 12 months following Pinatubo was 0.13 over Tsukuba, and the Pinatubo aerosol cooling was 3.1 Wm⁻². Recent stratospheric aerosol radiative cooling is about one-twentieth of that caused by the Pinatubo aerosols.





The surface temperature could be lowered by about 0.015–0.025 °C during the summer if we divide 0.3–0.5 °C by 20. It is very difficult to detect such a small change of surface temperature during one year. However, it is noteworthy that increased stratospheric aerosol radiative cooling continued for at least four years, from January 2008 to

- May 2012. Climate models have been used to simulate climate for a year after volcanic eruptions (Haywood et al., 2010; Kravitz et al., 2011), but multi-year simulations will be necessary to understand the effects of longer term increases in stratospheric aerosols, because, for example, the ocean integrates volcanic radiative cooling and responds over a wide range of time scales (Stenchikov et al., 2009).
- We next estimated the influence of the increase in stratospheric aerosols after volcanic eruptions on the XCO₂ determined by GOSAT. When the GOSAT XCO₂ is retrieved by using the 1.6-μm band without taking account of sulfuric acid particles in the stratosphere, the negative bias of XCO₂ is estimated to be 0.3 % (~ 1 ppm) for an AOT of 0.02 at 550 nm and surface albedo at 0.1 (Ota et al., 2008). It is noteworthy that the largest values of AOT at 532 nm after the volcanic eruptions of Mt. Sarychev and Mt. Nabro were equal to or larger than 0.02. A regional and time-dependent bias
- of 1 ppm is not small for surface CO₂ flux estimation (Rayer and O'Brien, 2001; Takagi et al., 2011). Therefore, it is necessary to take into account the effects of increased stratospheric aerosols for GOSAT XCO₂ retrieval (Uchino et al., 2012).

20 5 Concluding remarks

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An increase in stratospheric aerosols caused by the volcanic eruption of Mt. Nabro on 12 June 2011 was first observed by lidars at Tsukuba and Saga in Japan. The maximum backscattering ratios at 532 nm were 2.0 at 17.0 km on 10 July over Tsukuba and 3.6 at 18.2 km on 23 June over Saga. The maximum integrated backscattering coefficients above the first tropopause height to 33 km were $4.18 \times 10^{-4} \text{ sr}^{-1}$ on 11 February 2012 over Tsukuba and $4.19 \times 10^{-4} \text{ sr}^{-1}$ on 23 June 2011 over Saga.





Lidar observational results at Tsukuba from January 2008 through May 2012 revealed increases in stratospheric aerosols after the volcanic eruptions of Mt. Kasatochi in August 2008 and Mt. Sarychev Peak in June 2009. The yearly averaged IBCs at Tsukuba were $2.60 \times 10^{-4} \text{ sr}^{-1}$, $2.52 \times 10^{-4} \text{ sr}^{-1}$, $2.45 \times 10^{-4} \text{ sr}^{-1}$, and $2.20 \times 10^{-4} \text{ sr}^{-1}$ for 2008, 2009, 2010, and 2011, respectively. These values were about twice the IBC of 5 the background level $(1.21 \times 10^{-4} \text{ sr}^{-1})$ during the period from 1997 to 2001 at Tsukuba. The elevations of annual average AOT above background levels were about 0.0050-0.0070 from 2008 to 2011 based on an assumed lidar ratio of 50 sr. The negative radiative forcing (cooling) was then roughly 0.13–0.18 Wm⁻² for the same period based on a conversion factor of 25 Wm^{-2} from AOT to radiative forcing. These values are 10 not small compared to the radiative heating associated with increases in CO₂, about 0.28 Wm⁻² over the decade since 2000 (Solomon et al., 2011; NOAA, 2012). However, because the concentrations of these volcanic aerosols are not always spatially homogeneous, their radiative forcing might be overestimated. The influence of the increase in stratospheric aerosols caused by volcanic eruptions on GOSAT XCO₂ retrieval is

¹⁵ in stratospheric aerosols caused by volcanic eruptions on GOSAT XCO₂ retrieval non-negligible.

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Station Tsukuba Saga Transmitter Nd:YAG Nd:YAG Laser Wavelength 532 nm 1064 nm 532 nm 1064 nm Pulse energy 140 mJ 230 mJ 130 mJ 130 mJ Pulse repetition 20 Hz 10 Hz 0.2 mrad 0.2 mrad Beam divergence 0.2 mrad 0.2 mrad Receiver Telescope type Schmidt Cassegrain Schmidt Cassegrain 35.5 cm (Far) 30.5 cm Telescope Diameter 20.0 cm (Near) Field of view 1.0 mrad 1.0 mrad P and S Polarization P and S None None Number of 5 2 3 1 channels Vertical resolution 7.5 m (minimum) 7.5 m (minimum) PMT PMT Detectors APD APD (C30956EH) (R3234-01) (C30956EH) (R3234-01)

 Table 1. Characteristics of two-wavelength polarization lidar systems at Tsukuba and Saga.

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12 bit A/D and Photon Counting

12 bit A/D and Photon Counting

Signal processing



Fig. 1. Vertical profiles of the backscattering ratio *R* (pink line) and total depolarization ratio δ (%) (blue line) at $\lambda_2 = 532$ nm over Tsukuba (upper panel) and Saga (lower panel) from June 2011 through January 2012. Horizontal dashed lines show the first local tropopause heights. Large values of *R* and δ below tropopause heights are caused by cirrus clouds.





Fig. 2. Horizontal (upper panel) and vertical (lower panel) projections of isentropic forward trajectories of air parcels initially at an altitude of 17 km over Mt. Nabro (red square) versus time. The trajectories were calculated for ten days from 23:00 UT on 12 June 2011. Tsukuba and Saga lidar sites are indicated by red circles in the upper panel.







Fig. 3. Monthly means of geopotential height (m) and wind $(m s^{-1})$ on 100 hPa in June 2011 calculated from NCEP/NCAR reanalysis data. The wind speed scale is shown above the right side of the figure.





















Fig. 6. Profiles similar to Fig. 1 over Tsukuba from June to December 2009 after the 2009 Sarychev eruption.







Fig. 7. Temporal variation of the integrated backscattering coefficient (IBC) from the first tropopause to an altitude of 33 km (pink solid diamond) over Tsukuba from January 2008 through May 2012. The dotted line represents the seasonal variation of the monthly averaged IBC for background stratospheric aerosols observed at Tsukuba during October 1997 through September 2001. The date of each volcanic eruption is shown on the upper horizontal line.

