

Article

First In-Situ Measurements of Plume Chemistry at Mount Garet Volcano, Island of Gaua (Vanuatu)

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Abstract: Recent volcanic gas compilations have urged the need to expand in-situ plume measurements to poorly studied, remote volcanic regions. Despite being recognized as one of the main volcanic epicenters on the planet, the Vanuatu arc remains poorly characterized for its subaerial emissions and their chemical imprints. Here, we report on the first plume chemistry data for Mount Garet, on the island of Gaua, one of the few persistent volatile emitters along the Vanuatu arc. Data were collected with a multi-component gas analyzer system (multi-GAS) during a field campaign in December 2018. The average volcanic gas chemistry is characterized by mean molar CO₂/SO₂, H₂O/SO₂, H₂S/SO₂ and H₂/SO₂ ratios of 0.87, 47.2, 0.13 and 0.01, respectively. Molar proportions in the gas plume are estimated at 95.9 ± 11.6, 1.8 ± 0.5, 2.0 ± 0.01, 0.26 ± 0.02 and 0.06 ± 0.01, for H₂O, CO₂, SO₂, H₂S and H₂. Using the satellite-based 10-year (2005–2015) averaged SO₂ flux of ~434 t d⁻¹ for Mt. Garet, we estimate a total volatile output of about 6482 t d⁻¹ (CO₂ ~259 t d⁻¹; H₂O ~5758 t d⁻¹; H₂S ~30 t d⁻¹; H₂ ~0.5 t d⁻¹). This may be representative of a quiescent, yet persistent degassing period at Mt. Garet; whilst, as indicated by SO₂ flux reports for the 2009–2010 unrest, emissions can be much higher during eruptive episodes. Our estimated emission rates and gas composition for Mount Garet provide insightful information on volcanic gas signatures in the northernmost part of the Vanuatu Arc Segment. The apparent CO₂-poor signature of high-temperature plume degassing at Mount Garet raises questions on the nature of sediments being subducted in this region of the arc and the possible role of the slab as the source of subaerial CO₂. In order to better address the dynamics of along-arc volatile recycling, more volcanic gas surveys are needed focusing on northern Vanuatu volcanoes.

Keywords: Vanuatu; Gaua; Mount Garet; Multi-GAS; volcanic gas compositions; volatile fluxes

1. Introduction

Monitoring the composition and mass flux of subaerial volcanic emissions is key to understanding how volcanism impacts our planet, on both local and global scales.

At subduction zones, de-volatilization of subducting slab materials during high-pressure metamorphism [1–3] returns volatiles back to the hydrosphere-atmosphere via mantle wedge melting and arc volcanism. These materials are injected into the mantle through subducted sediments [4–7], altered oceanic crust (AOC) [8,9] and serpentinized lithospheric mantle [10–12].

Therefore, constraining major volatile budgets from subaerial volcanic arc emitters provides a tool to partially quantify the recycling efficiency of H₂O, CO₂, SO₂ and other volatiles via the subduction process. The contrasting volatile ratios in the depleted mantle (CO₂/S_T of ~0.3–0.8) [13], subducted sediments (CO₂/S_T ~1–100) [14] and crustal fluids from limestone de-carbonation (CO₂/S_T >>100) [15] imply that measuring volcanic gas composition can bear crucial information on volatile sources at depth. At the arc-scale, ever-increasing volcanic gas inventories are key to understanding the volatile cycle at subduction.

However, global volcanic gas datasets [15–22] are still sparse and incomplete, largely due to the remote location of some volcanic centers, inaccessibility, social/political instability of geographic regions and/or high level of volcanic activity.

For instance, the Southwestern Pacific region is home to some of the most persistently degassing volcanoes on Earth [23], but Melanesian volcanoes have systematically been underrepresented in previous literature reviews. Recent advances in satellite remote sensing techniques, such as the ozone monitoring instrument (OMI), have greatly improved our knowledge on SO₂ flux emissions globally [23], revealing Vanuatu as a major volcanic SO₂ source globally, and urging the volcanic gas community to target volcanic gas studies in the poorly studied, yet strongly emitting volcanoes along this arc segment.

In-situ measurements of volcanic gas compositions can be attained by using relatively inexpensive multi-component gas analyzer systems (such as the multi-GAS) [24,25]. In addition to their great potential in eruption forecasting, these portable units have revolutionized our understanding of along-arc chemistry of volcanic gases at different regions of the world. Moreover, when combined with remotely sensed SO₂ fluxes, these measurements provide a means to constrain volcanic gas fluxes, especially CO₂ [17–21,26–29].

In the Vanuatu arc, volcanic gas surveys have mostly focused on Yasur [30–32] and Ambrym [33–35] due to the persistent high levels of degassing and volcanic activity. However, Reference [36] reported the first estimate of volcanic SO₂ budget for the Vanuatu Island Arc, in which the authors highlight other vigorously degassing volcanoes along the arc, for which volcanic gas compositions have remained unconstrained.

In this paper, we present the results of a field campaign to measure gas compositions and estimate total volatile fluxes emitted from Mount Garet volcano (or Mount Gharat; also occasionally referred to in the literature as Gaua volcano), located on the island of Gaua, Vanuatu, whose activity in 2009–2010 prompted the evacuation of more than 1500 inhabitants [36]. Data collected in December 2018 is here integrated with previous SO₂ flux reports from both ground-based measurements [36] and satellite records [23] in order to assess the volatile budget of Mount Garet. Data from this study represents the first in-situ plume composition measurements obtained for Mount Garet and thus provides information on volcanic gas signatures in the northern part of the arc segment. We also integrate our novel dataset in the context of previously reported gas compositions for other Vanuatu volcanic centers and discuss the overall implications of this and other recent studies of Melanesian volcanoes [37] to global volatile budgets.

2. The Vanuatu Island Arc

The New Hebrides Island Arc is located in the southwest Pacific (Figure 1). It is composed of a narrow chain of Neogene to recent volcanic islands originating from the eastward subduction of the Australian plate beneath the Pacific plate, with associated back-arc extension of the North Fiji basin on the Pacific side [38,39]. The recent tectonic evolution of the arc is dominated by its collision with the D'Entrecasteaux Zone (DEZ) [40], which began 1.5–3 Ma ago near Epi island, and has been migrating

northward as a result of the obliquity of the DEZ relative to the trench [40,41]. The New Hebrides Island Arc is subdivided in four different segments, the northernmost being essentially submarine and poorly studied. The other three include the central and south segments (CVS and SVS, respectively) and the high-Mg andesite volcanic segment (HMAVS) [42].

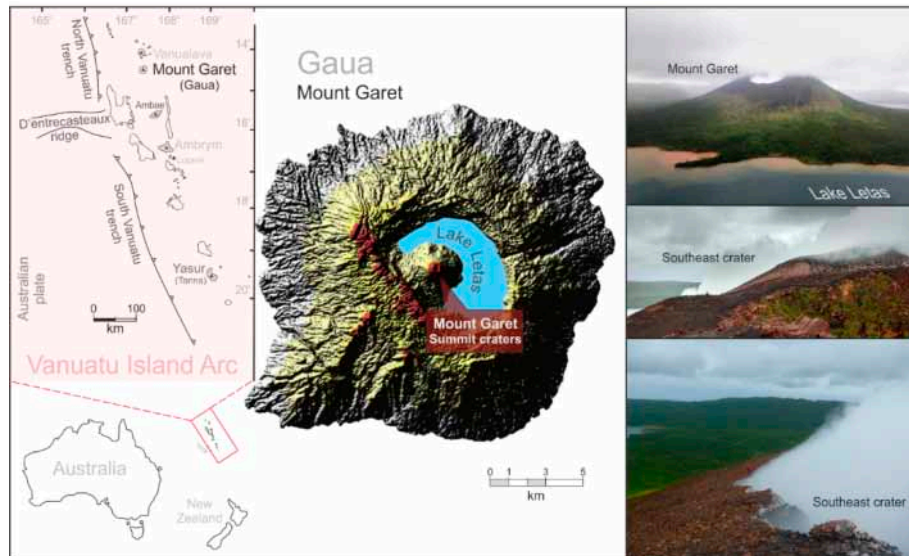


Figure 1. On the left: illustration of the overall tectonic context of the Vanuatu Island Arc, with currently active volcanoes and their location marked in red triangles. In the center: digital elevation model (DEM) of the island of Gaua; land features above 700 m (a.s.l.), including Mount Gare's summit craters, are highlighted in red. On the right: photos of Mount Gare and its currently degassing southeast crater (December 2019); photos of National Geographic property.

Vanuatu extends over the CVS and SVC, for about 700 km. It consists of a partly emergent ridge with an average width of 200 km, that overlies a steeply eastward-dipping subducted slab [42]. The DEZ collision triggered the detachment of the Australian plate lithosphere beneath the central part of Vanuatu. However, this detachment is less pronounced in the Northern part of the Vanuatu arc (due to a westward and upward invasion of the Indian-type MORB mantle) [42] where the island of Gaua is located.

Gaua, also named Santa Maria, represents the emerged part of a 3000 m-high and 40 km-wide composite volcano [43]. It consists of a little-dissected symmetrical cone, about 25 km-across, truncated at an elevation between 500 m and 690 m (a.s.l.) by an 8.5×6 km central caldera, filled by a large crater lake (Lake Letas; Figure 1). The active cone of Mt. Gare rises in the middle of Lake Letas.

Mt. Gare (797 m, a.s.l.) has two distinct craters. Since 1991, the activity has clustered on the southeast crater, which was also the site of the 2009–2010 explosions [44] and the survey location for the in-situ volcanic gas measurements reported here. A previous study [45] on the geology of the volcano identified a thick pile of basaltic and andesitic lava flows with associated rubble horizons and beds of red cinders or agglomerates. Mallick and Ash [45] also recognized an extensive cap of ash surrounding the caldera. On the flanks, small parasitic cones and fissure-fed lava flows are distributed along fractures concentric to the caldera rim. The morphology of vents and lava surfaces mantling the lower slopes suggest that these peripheral extrusions are very young. Analysis of 2009–2010 lavas, products of the last major eruption, revealed identical compositions to those previously reported in literature [43,45,46]. The occurrence of basaltic andesite and basaltic trachyandesite volcanites with highly calcic plagioclase suggest that the 2009–2010 eruption of Gaua was fed by an alkaline (K-rich) magmatic source, typical of the northern part of the Vanuatu arc [44].

The 2009–2010 Eruption of Mount Garet (Gaua)

According to local inhabitants, the 2009–2010 eruption of Mount Garet was the strongest witnessed since 1963. Bani et al. [44] assessed magma composition, lithology of volcanites, ash mineralogy, SO₂ mass discharge, halogen release and lake water chemistry, and provided a comprehensive understanding of the 2009–2010 sequence of eruptive events described below.

A series of mild explosions on 29 September 2009, marked the beginning of a year-long eruption at Mount Garet. Shortly after, on October 3 and 14, airborne differential optical absorption spectroscopy (DOAS) measurements revealed SO₂ fluxes of 3024 ± 1063 and 2765 ± 956 t d⁻¹, respectively [36]. Despite high degassing rates, the first two months of volcanic activity were largely characterized by low explosivity, with the exception of four large explosive events that took place between 18–27 November 2009 and that displaced 250 inhabitants from the western coast of Gaua, the area most affected by ash fallout. By 17 December, SO₂ discharge rates had remained high (3084 ± 1080 t d⁻¹) [36], although the low proportions of juvenile materials and the abundance of hydrothermally altered rock and lava fragments deposited until then suggested an initial eruptive phase fueled by the subsurface interaction between the rising magma and a hydrothermal system [44].

Volcanic activity intensified again in January 2010. Eruption columns regularly reached up to 3000 m a.s.l. over the following five months (Volcanic Ash Advisory Center, Darwin) (GVP: 05/2011 (BGVN 36:05)) and ash fallout extended at least 10 km away from the source [44]. Following this, magma composition became more basic and more homogeneous, as evidenced by the lithological characteristics of ash fragments and further by Scanning Electron Microscopy (SEM) analysis [44]. The prominent role of more juvenile magma in this eruptive phase is attested by larger proportions of vesiculated fragments, and by a concomitant decrease in hydrothermally altered materials emitted. By the end of January, up to 75% of the erupted products were juvenile, and composed of 50% of vesiculated fragments [44].

Finally, in July 2010, volcanic activity at Mount Garet started to decline and the eruption ended in September 2010. The Vanuatu Geohazards Observatory (VGO) maintained the alert level at two throughout the eruption, and no further evacuation was carried out. During the 2009–2010 Gaua eruption, Lake Letas also went through a partial color change from blue to brown [44], which could still be seen in December 2018 in the area adjacent to Mt. Garet (Figure 1).

3. Materials and Methods

We used a multi-GAS unit [24,25] to measure in-plume concentrations of major volatile species at Mount Garet's southeast crater (~797 m a.s.l.). The instrument was composed of a Gascard nondispersive infrared (NDIR) CO₂ spectrometer from Edinburgh Sensors and City Technology SO₂, H₂S and H₂ electrochemical sensors (see supplementary material S1 for details on sensor's manufacture models, sensor's accuracy and repeatability, calibration ranges and gas standards). Gas sensors were calibrated before and after the field campaign at the Earth and Marine Sciences Department (DiSTeM) of the University of Palermo. The system also included temperature (T) and relative humidity (Rh) KVM3/5 Galltec-Mela sensors. All components and respective power sources were contained inside a weather-resistant plastic case, with inlet/outlet ports to provide access to ambient air.

In the field, volcanic gas was pumped through the in-series connected sensors using a small pump with a flow rate of 1.8 l/min. In-plume gas concentrations were measured non-continuously for approximately 4 h (see Figure 2 for the whole concentration time series of CO₂ and SO₂) at a frequency of 1 Hz. Each measurement routine was preceded by instrument warm-up (3 min) and two minutes of ambient air flow to flush residual volcanic gas that remained trapped within the circuit.

Multi-GAS data were processed using the Ratiocalc software [47]. We selected specific acquisition time windows showing good temporal correlation ($R^2 \geq 0.70$) between volatile concentrations measured simultaneously [48–50] (see Figure 3a for different gas species). Molar ratios and errors reported in Table 1 were calculated from the slope and the R^2 value of the best-fit line (linear regression, such as the example of Figure 3b, in black), respectively. Following an identical procedure, bulk compositions

and associated errors reported in Tables 1 and 2 were obtained from all data points for a given x/SO_2 pair. The x/SO_2 dot-to-dot ratio versus in-plume SO_2 concentration [47,51] (Figure 3a) is also shown to demonstrate that ratios, widely scattered at low SO_2 concentrations, converge to a less-varying ratio value (statistically indistinguishable from bulk x/SO_2 ratios reported in Table 1) at high SO_2 concentrations (30–70 ppmv).

Table 1. Individual x/SO_2 molar ratios measured by the multi-component gas analyzer system (multi-GAS) in the Mount Garet’s plume. These correspond to time windows showing good temporal correlation between gas species ($R^2 \geq 0.70$) and strong volcanic gas signal (≥ 30 ppm of SO_2). Ratios and associated errors correspond to the slope and the R^2 value of the best-fit regression line, respectively.

Gas Species	Molar Ratio x/SO_2	R^2	Error (\pm)	SO_2 Max (Plume Marker; in ppm)	Nr. of Samples
H_2	0.04	0.73	0.006	36.1	713
	0.04	0.70	0.007	39.9	804
	0.03	0.86	0.009	56.8	288
	0.03	0.78	0.005	61.9	534
	0.03	0.74	0.007	56.3	480
Bulk composition \pm	0.03 0.006				
H_2O	40.5	0.70	7.4	63.2	431
	40.0	0.73	5.7	43.8	1006
	47.8	0.78	6.3	61.9	1387
	55.5	0.86	3.7	39.9	544
	53.2	0.71	4.7	39.9	1538
Bulk composition \pm	47.2 5.7				
CO_2	0.85	0.82	0.076	36.6	557
	1.07	0.71	0.182	37.8	473
	0.65	0.87	0.045	50.9	451
	1.02	0.76	0.265	31.8	439
	1.06	0.70	0.653	37.3	725
Bulk composition \pm	0.87 0.24				
H_2S	0.13	0.93	0.011	52.9	526
$\text{H}_2\text{O}/\text{CO}_2$	68.6	0.75	10.0	61.9	1019

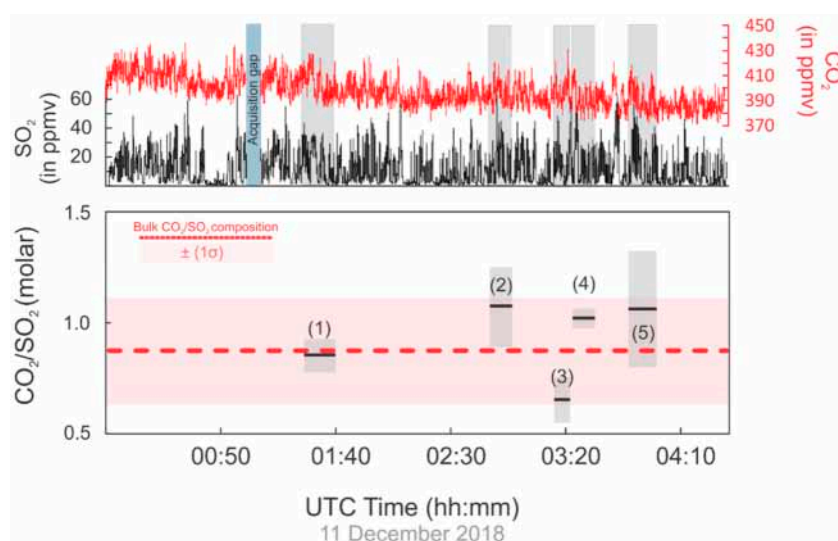


Figure 2. CO_2/SO_2 mixing ratios estimated across the 4-h survey period. At the bottom, the dash red line represents the regression analysis of all 5 CO_2/SO_2 ratios (Table 1) combined. SO_2 and CO_2 concentration (in ppmv) time series at the top illustrates plume conditions at which individual ratios were calculated. Note that CO_2/SO_2 ratio (3) is then shown in Figure 3 for detailed calculation procedure.

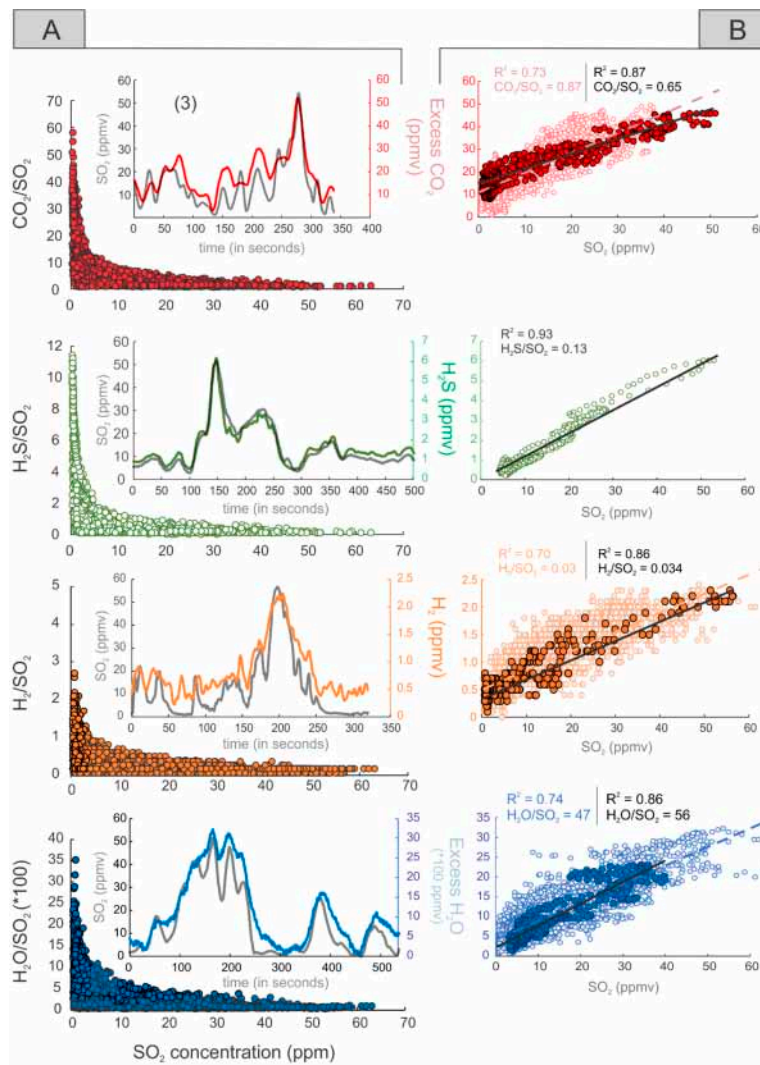


Figure 3. x/SO_2 Mixing ratios in the Mount Garet plume. (A): scatterplot of dot-to-dot x/SO_2 molar ratios versus SO_2 concentrations, showing that the ratios converge to a single value at high SO_2 concentration (Tamburello, 2015); and concentration (ppmv) time series of each species (CO_2 in red, H_2S in green, H_2 in orange and H_2O in blue) with SO_2 (in black) in the Mount Garet plume (note that the x -axis gives the duration of each survey in seconds). (B): Least-square regression analysis are shown for each species versus SO_2 ; Slope and R^2 regression values in black correspond to the specific time window shown in A; colored values correspond to the regression analysis of bulk data reported in Table 1.

Table 2. Bulk composition of Mount Garet’s plume; x/SO_2 ratios correspond to the gradients of the best-fit regression lines of all combined ratios reported in Table 1 for a given x/SO_2 pair. Errors are expressed as the standard error of the regression analysis and subsequent error propagation.

Gas Species	Molar Ratio x/SO_2	$\pm (1\sigma)$	%Err	Composition (mol%)	$\pm (1\sigma)$
H_2O	47.2	5.7	12	95.88	11.58
CO_2	0.87	0.24	24	1.77	0.49
SO_2	1.00	—	—	2.03	—
H_2S	0.13	0.01	8	0.26	0.02
H_2	0.03	0.01	20	0.06	0.01

Cross-sensitivity SO_2 effects on the H_2S sensor were estimated at 14% during calibration and corrections were applied during data processing to calculate interference-corrected H_2S concentrations in the gas plume.

Water concentration ratios were calculated using measurements of T, P and Rh and the Arden Buck equation [47]. CO_2 and H_2O concentration ratios were estimated after subtraction of background-air concentration ratios acquired at plume-free areas, and their relative abundance is given in Figure 4 following the above described procedure.

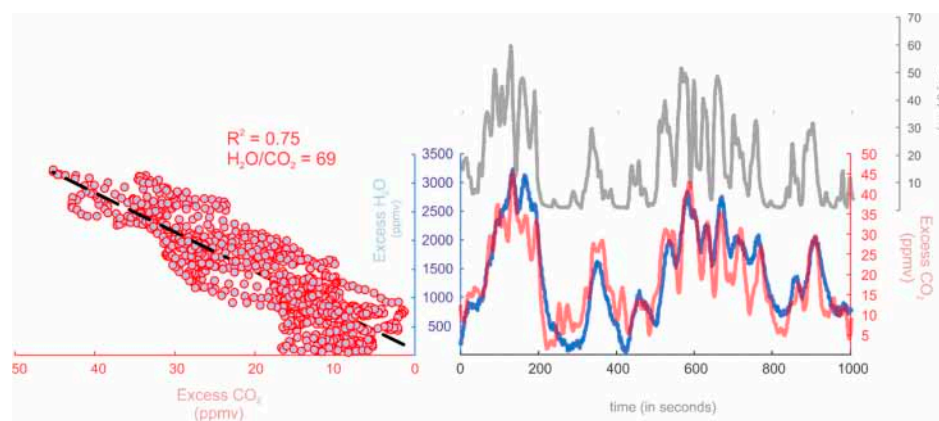


Figure 4. On the left: scatter plot of H_2O vs. CO_2 mixing ratio in the Mount Garet plume; the least-square regression line is shown as black dashed line on each plot ($R^2 = 0.75$). On the right: concentration (ppmv) time series of H_2O (blue) and CO_2 (red) recorded at a frequency of 1 Hz; in black, SO_2 concentration (ppmv) is given as an analog for plume density during our measurement period.

4. Results

Volcanic Gas Composition

Gas emissions at Mount Garet were measured at close proximity to the southeast crater. Therefore, over the non-continuous, 4-h measurement period, the multi-GAS consistently captured a strong plume signal (up to ~ 70 ppm SO_2). Background air-corrected CO_2 concentrations varied from <1 to ~ 60 ppm (Figures 3 and 4). Rare plume batches yielded H_2S concentrations above the 14% cross-sensitivity to SO_2 but occasionally reached in-plume concentrations up to ~ 7 ppm (Figure 3). We recorded H_2 concentrations up to 3 ppm (Figure 3). The volcanic H_2O signal (up to ~ 3500 ppm; Figures 3 and 4) was intermittently resolvable from background noise caused by high humidity conditions characteristic of tropical environments.

Derived volatile ratios (in the form of x/SO_2 ratios; where x is either H_2O , CO_2 , H_2S , or H_2) for Mount Garet were obtained in measurement windows where strong positive co-variations ($R^2 \geq 0.70$) were observed between SO_2 and the other volatile species measured (Figure 3). Interference-corrected $\text{H}_2\text{S}/\text{SO}_2$ ratio and H_2/SO_2 ratios (0.13 ± 0.01 and 0.03 ± 0.01 , respectively) remained constant throughout the entire measurement period. Likewise, CO_2/SO_2 ratios varied narrowly between 0.65 and 1.07 (avg. 0.87 ± 0.24). Due to variable mixing between atmospheric H_2O and the magmatic plume, $\text{H}_2\text{O}/\text{SO}_2$ measurements were more variable, ranging from ~ 40 to ~ 56 (avg. 47.2 ± 5.7 ; Tables 1 and 2). Similarly, $\text{H}_2\text{O}/\text{CO}_2$ ratios also showed wide variations across the entire 4-h window, reaching values as low as ~ 40 and as high as ~ 70 . However, the best correlation ($R^2 = 0.75$) was found at the upper limit of that range, yielding a $\text{H}_2\text{O}/\text{CO}_2$ ratio of ~ 69 (Figure 4). The volcanic plume composition of Mount Garet was estimated at 95.9 mol% H_2O , 1.8 mol% CO_2 , 2.0 mol% SO_2 , 0.3 mol% H_2S , and 0.06 mol% H_2 (uncertainties reported in Table 2 are expressed as the standard error of the regression analysis and subsequent error propagation).

5. Discussion

Our observations constrain the gas plume composition at Gaua as hydrous (~95.9 mol% H₂O), with SO₂ dominating over H₂S (H₂S/SO₂ of 0.13 ± 0.01), and with a H₂/H₂O ratio of ~0.001.

Therefore, by using gas-phase H₂/H₂O (~0.001) and H₂S/SO₂ (~8.4) redox couples and using published thermodynamic constants [52,53], we can estimate the oxygen fugacity (fO₂) of Mount Garet’s volcanic gases (e.g., Moussallam et al., 2019). Equations (1) and (2) also provide the means to estimate the gas-melt equilibrium temperature:

$$\log \frac{H_2}{H_2O} = \frac{-12707}{T} + 2.548 - \frac{1}{2} \log fO_2 \tag{1}$$

$$\log \frac{SO_2}{H_2S} = \frac{27377}{T} + \frac{3}{2} \log fO_2 - \log fH_2O \tag{2}$$

The value of *f*H₂O used in (2) is 0.959 given that, at ~1 bar the (P_{tot} × nH₂O)/n_{tot} = [(1 bar) (0.959n_{tot})]/n_{tot} = 0.959 bar (where fugacity of a gas is equal to its partial pressure and that P(H₂O) = P is the pressure in bar and n_i the amount of species *i* in mol%) [54].

Solution yields an equilibrium temperature of ~641 °C and a log*f*O₂ equivalent to ΔQFM = +2.0 where QFM refers to the quartz-fayalite-magnetite buffer [55], and where ΔQFM = log*f*O₂ – log*f*O₂ of QFM at the corresponding temperature range estimated above. This solution plots well within the global trend highlighted in [22].

As such, an equilibrium temperature of ~641 °C is higher than the temperature at which sulfur partitioning into the liquid hydrothermal system or gas scrubbing reactions in a shallow subsurface environment may occur [56,57]. Therefore, the absence of contamination of Gaua’s magmatic gas allows for comparison with other high-temperature arc gases worldwide (Figure 5) [17,22,28]. Being the first for Gaua, our results contribute to our understating of volcanic gas signature and fluxes along the Vanuatu arc, as discussed below.

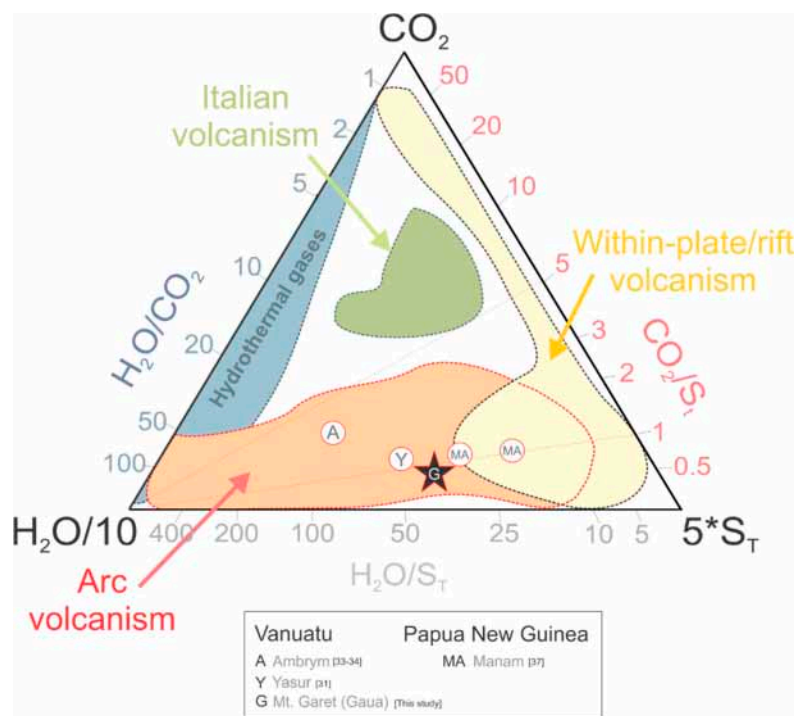


Figure 5. H₂O–CO₂–S_T (S_T = SO₂ + H₂S) ternary plot with gas composition for Mount Garet volcano (in black). Ambrym [33,34] and Yasur [31] are also plotted for comparison, as well as Manam (Papua New Guinea) [37]. For within-plot field delimitations (e.g., hydrothermal gases and arc volcanism) see [28].

5.1. Along-Arc and Inter-Arc Volcanic Gas $\text{CO}_2/\text{S}_\text{T}$ Variations

Recent global volcanic gas compilations have revealed systematic inter-arc and along-arc variations in gas $\text{CO}_2/\text{S}_\text{T}$ ratios [15,21] which are thought to be mainly governed by variations in mineralogy and chemistry of sediments (especially their C load) subducted at the corresponding trenches [6,7]. A better understanding of these global correlations between arc inputs (sediments) and outputs (volcanic gases) requires extending volcanic gas measurements to remote, poorly studied arc regions, such as Melanesia. Indeed, as noted by Aiuppa et al. [15,21] several of the top volcanic arc emitters have yet to be measured for their $\text{CO}_2/\text{S}_\text{T}$ ratios, due to their remote location and/or high level of volcanic activity.

The results we report here add new data to this global dataset to allow a more complete characterization of the source, fate and cycling of volatiles, especially C, in subduction zones. In this section, we focus the discussion of our results within the context of recently collected gas data for other Melanesian volcanoes.

As two of the most active volcanoes in the world, volcanic gas emissions at Yasur and Ambrym (Vanuatu; see Figure 1) have been studied extensively in recent years. First reports on volcanic plume chemistry at Ambrym were presented by Allard et al. [33,34], while Oppenheimer et al. [30], Métrich et al. [31] and Woitischek et al. [32] reported on gas compositions at Yasur volcano. These studies inferred the plume volcanic gas $\text{CO}_2/\text{S}_\text{T}$ ratios at ~ 1.5 [15,33,34] and ~ 1.6 [31] at Ambrym and Yasur, respectively, within the range of CO_2 -poor (Group 1) volcanic gases ($\text{CO}_2/\text{S}_\text{T} < 2$) [15,21].

Aiuppa et al. [15,21] noted that this CO_2 -poor volcanic gas composition for the Vanuatu arc, as inferred from observations at Ambrym and Yasur, is somewhat at odds with the relatively high C content of sediment in the subducting slab (bulk sediments in the Vanuatu trench contain up to 4 wt.% CO_2) [6]. In other arc sectors where C is subducted via sediments (e.g., Central and northern Southern America) [7], volcanic gases are typically more C-rich, and display characteristics of those volcanoes in the Group 2 category ($2 < \text{CO}_2/\text{S}_\text{T} < 4$) [15,21,50,58]. In contrast, the Vanuatu gas compositions measured thus far resemble more closely those of subduction zones of the NW Pacific (the Kuril–Kamchatka and Marianas–Japan arc segments) [15], all of which exhibit low C content in subducting sediments [6].

Our results here provide a first insight into the $\text{CO}_2/\text{S}_\text{T}$ signature of volcanic gases in the northern part of the Vanuatu arc. Our average volcanic gas $\text{CO}_2/\text{S}_\text{T}$ estimate at Gaua is ~ 0.9 , for measurements performed in very dense plume conditions (up to >50 ppmv SO_2). The 2018 Gaua gas composition also plots at the low- CO_2 end of the population of volcanic arc gases worldwide (Figure 5). Notably, a similarly C-poor gas measurement was recently obtained further to the north at Manam volcano, in Papua New Guinea, for which Liu et al. [37] quoted a $\text{CO}_2/\text{S}_\text{T}$ ratio of ~ 1 .

The relatively high C content of sediments at the Vanuatu trench led Aiuppa et al., [21] to predict a $\text{CO}_2/\text{S}_\text{T}$ ratio of ~ 2.5 for Gaua, well in excess of our 2018 measured ratio of ~ 0.9 . Thus, the C-rich nature of the subducted sedimentary input in the region does not appear to translate into a C-rich volcanic gas output, as seen elsewhere (e.g., central and northern Southern America). The possibility that the sedimentary C input into the subduction zone is accreted to the accretionary wedge (rather than being subducted) is inconsistent with the results of Von Huene and Scholl [59], who classified the Vanuatu arc as a non-accreting margin, with effectively no accretionary prism present. Further, seismic reflection data indicate that most of the sediments are indeed subducted [60]. In fact, the same authors argue that, in the region, the addition of a significant subducted sediment component has variably modified the mantle wedge over time. Within the less-surveyed northern Vanuatu region, distinct geochemical patterns can be observed. For instance, lavas from Gaua have lower $^{143}\text{Nd}/^{144}\text{Nd}$ than lavas from Merelava, despite similar Sr- and Pb-isotope compositions. This can be explained in terms of a common Indian-MORB-like mantle source but would indeed require a higher contribution from subducted sediments to Gaua lavas [60]. Ultimately, the mismatch between the high C input (sediments) and the low C output (volcanic gases) may simply reflect a limited C recycling by fluids delivered by the slab, leading to sedimentary C transport deep in the mantle [7].

We caution, however, that our discussion is based on one single, brief campaign at Gaua, while more continuous and prolonged gas records would be required to fully capture the volcano's time-averaged

degassing behavior and gas composition. Indeed, gas compositions exhibit both long-term changes in response to changes in volcano behavior (e.g., as persistent degassing activity transitions into unrest and finally eruption) [19], and also more subtle short-term changes related to rapid events of gas segregation and explosive burst. At Yasur volcano, for example, the bulk composition appears to have remained relatively stable since the 2000s, yet with CO_2/S_T ratios spanning between 1.6 and 2.2 [30–32]. In addition, short-term (timescales of seconds to minutes) fluctuations in gas compositions between ‘passive’ ($\text{CO}_2/\text{S}_T \sim 1.9$) and ‘active’ ($\text{CO}_2/\text{S}_T \sim 2.9$; during strombolian explosions) degassing regimes have also been observed [30,32]. Ultimately, additional gas observations at Gaua and other Vanuatu volcanoes is required for a more robust assessment of C recycling efficiency along the arc.

5.2. Implications for the Vanuatu Arc Volatile Budget

Vanuatu hosts two of the largest volcanic gas emission sources globally. Ambrym, a volcanic island known for frequently hosting the lava lakes of Benbow and Marum until its 2018 eruption [61] appears first on the list of subaerial SO_2 emissions measured from space between 2005–2015 [23]. The 10-year averaged SO_2 emission inventory ranks Yasur volcano (located in the island of Tanna; see Figure 1) at number 11 in the list. Notably, Bagana, Tavurvur and Manam, from the nearby archipelago of Papua New Guinea, are all in the top 10. Clearly, the southwest Pacific is a major contributor to global volcanic gas release.

Vigorous volcanic SO_2 degassing in Vanuatu had been reported previously by Bani et al. [36]. Using repeated DOAS (differential optical absorption spectroscopy) ground measurements (2004–2009) and satellite OMI data (2004–2011), (Bani et al., 2012) estimated that on average 3 Tg y^{-1} of volcanic SO_2 are emitted into the atmosphere along the Vanuatu arc, to which Ambrym reportedly contributed to about a third. Such estimates did not include sporadic eruptions and/or extreme passive degassing events, but nonetheless represented about one-fifth of currently estimated global volcanic SO_2 emissions. The study also marked the first ground-based SO_2 flux estimates for Mount Garet volcano. During October and December 2009, seven DOAS traverses yielded a mean SO_2 emission rate of $\sim 2959 \text{ t d}^{-1}$ [36]. These measurements were taken during the 2009–2010 eruption that started in September (see cfr. 2 for details) and most probably reflected the high degassing rates registered in the first few months of the eruption [36,44].

In fact, Carn et al. [23] found much lower SO_2 emission rates for the period of 2005–2015 at Gaua. They reported a 10-year maximum of $\sim 1418 \text{ t d}^{-1}$ in 2012, and a 10-year average of $\sim 434 \text{ t d}^{-1}$ of SO_2 . Mismatches between ground and satellite-based emission inventories may arise from several factors [62]. Especially during the course of the 2009–2010 eruption (OMI satellite data yielded yearly averaged emissions of ~ 490 and 605 t d^{-1} for 2009 and 2010, respectively) [23], discrepancies may originate from difficulties in accurately measuring SO_2 in proximal ash-laden plumes, like the ones originated, for instance, in the explosive events of November 2009 [44]. Due to the low altitude of Mount Garet ($\sim 797 \text{ m a.s.l.}$), it is also possible for some passive degassing emission rates measured via satellite to be underestimated. That is because the detection limit will be lowest for low-latitude volcanoes, which benefit from more satellite observations under optimal conditions (e.g., low solar zenith angles) [23,62].

No SO_2 flux record was obtained during our 2018 survey. We therefore combine the 10-year (2005–2015) average volcanic SO_2 flux of Carn et al. [23] with our in-situ measurements of volcanic gas compositions (2018) to estimate, for the first time, Mount Garet’s emission inventory for H_2O , CO_2 , H_2S and H_2 . In doing so, we assume the 2005–2015 average [23] as the available best proxy for the characteristic SO_2 emission behavior during the volcano’s persistent (yet quiescent) degassing behavior (as observed during our in-situ survey in 2018). Using Bani et al. [36] SO_2 flux dataset would have led to substantially higher fluxes, yet most likely biased due to the fact that the large pulses in SO_2 release in the Bani et al. [44] dataset correspond to degassing behaviors associated with the 2009–2010 explosive eruptions and are markedly distinct from the passive degassing observed in 2018.

Our estimated fluxes are thus $\text{H}_2\text{O} \sim 5758 \text{ t d}^{-1}$, $\text{CO}_2 \sim 259 \text{ t d}^{-1}$, $\text{H}_2\text{S} \sim 30 \text{ t d}^{-1}$, and $\text{H}_2 \sim 0.5 \text{ t d}^{-1}$, and correspond to a total volatile flux (TVF) of about 6482 t d^{-1} for Mount Garet.

Our inferred CO_2 fluxes are especially relevant because they contribute new data to recent cataloguing efforts [19–21,29] aimed at a more robust quantification of the global volcanic CO_2 output. Such global estimates have remained poorly constrained until recently, because technical challenges in volcanic CO_2 measurements have resulted in sparse and incomplete global CO_2 flux catalogues. With a CO_2 flux of $\sim 259 \text{ t d}^{-1}$, Mount Garet ranks as intermediate-size volcanic CO_2 source (the median volcanic CO_2 flux in catalogues [20,29] is $\sim 320 \text{ t d}^{-1}$). Notably, our measured Gaua CO_2 flux is a factor ~ 3 lower than the CO_2 flux ($\sim 745 \text{ t d}^{-1}$) predicted by Aiuppa et al. [21]. The latter authors circumvented the lack of direct gas measurements at Gaua (unavailable at the time of their study) by predicting the time-averaged volcanic gas $\text{CO}_2/\text{S}_\text{T}$ ratio from trace element compositions (Ba/La and Sr/Nd ratio) of Gaua lavas and global gas vs. trace element relationships. The mismatch between the direct (this study) and indirect [21] estimates reflect the difference between measured and predicted $\text{CO}_2/\text{S}_\text{T}$ ratio (0.9 vs. 2.5), as discussed above (cfr. 5.1).

Our results suggest persistent degassing at Mt Garet (Gaua) contributes an additional $\sim 2\%$ to CO_2 emissions along the Vanuatu arc, which during the 2005–2015 period appeared dominated by prodigious lava lake degassing at Ambrym [33,34] (the inferred 2005–2015 average is $\sim 7586 \text{ t d}^{-1}$ [21]) and, to a lesser extent, by open-vent degassing at Aoba and Yasur (inferred 2005–2015 averages of ~ 4933 and $\sim 1549 \text{ t d}^{-1}$, respectively) [21].

6. Conclusions

Global inventories of volcanic gas data remain sparse and incomplete, especially for remotely located volcanic centers such as Mount Garet, in Gaua. Less than 60 of the several hundred currently degassing Holocene volcanoes have been characterized geochemically for their gas emissions and less than a few (<10) volcanoes have permanent in-situ instrumentation operating in real-time.

In this study we presented volcanic gas composition data from Mount Garet volcano. Our survey provides the first characterization of plume chemistry for this volcano. Although results reported here are punctual, they set unprecedented information on the composition and flux of volcanic gases at Mount Garet during a quiescent activity period (December 2018). By combining our results with satellite-based SO_2 flux measurements (average of 434 t d^{-1} over the 2005–2015 period) we estimated H_2O , CO_2 , H_2S , and H_2 outputs of $\sim 5758 \text{ t d}^{-1}$, $\sim 259 \text{ t d}^{-1}$, $\sim 30 \text{ t d}^{-1}$, and $\sim 0.5 \text{ t d}^{-1}$. These results rank Mount Garet as an intermediate-size volcanic CO_2 point source, well below the intense CO_2 emissions derived for nearby Ambrym (until its 2018 eruption) and Yasur volcanoes further south in the same Vanuatu arc. We caution that our observations are only likely to be representative of Mt. Garet's quiescent emissions, while much higher CO_2 and total volatile fluxes are likely to be associated with eruptive phases, as for instance supported by the high SO_2 flux emissions measured during the 2009–2010 period of unrest (for which no gas composition is unfortunately available). Our results thus confirm the need to better constrain the temporal variability of gas compositions through periodic and/or continuous surveys. Resolving the volatile output for both quiescent and unrest phases, along with newly reported volcanic gas plume compositions, would foster the ability to interpret volcanic gas monitoring signals and perform hazard assessments in the future.

Our volcanic gas survey has also revealed an especially C-poor composition ($\text{CO}_2/\text{S}_\text{T} \sim 0.9$) for Mount Garet volcano, well below the compositions derived for nearby Ambrym and Yasur ($\text{CO}_2/\text{S}_\text{T} \sim 1.5$ to 1.6), and un-matching with the previously predicted volcanic gas compositions for the same volcano ($\text{CO}_2/\text{S}_\text{T} \sim 2.5$). The contrast between the relatively high C content of sediments at the Vanuatu trench and the CO_2 -poor nature of volcanic gases measured at Mount Garet during our 2018 field campaign is intriguing. In other volcanic arcs where C is subducted through sediments (northern segments of the Andean Volcanic Belt, for example), volcanic gases typically display higher $\text{CO}_2/\text{S}_\text{T}$ ratios (~ 2 –4). Furthermore, despite the fact that recent gas studies at other persistently degassing Melanesian volcanoes have reported similar in-plume volatile signatures, more volcanic gas surveys are needed to

accurately assess the apparent mismatch between the high C input (sediments) and the low C output (volcanic gases). That will ultimately allow for more robust arc-scale interpretations of the nature of gas emissions from Southwest Pacific volcanism, a region of much relevance to arc CO₂ budgets globally, and their implications for our overall understanding of global geochemical cycles and atmospheric evolution over time.

Supplementary Materials: The following are available online at <http://www.mdpi.com/2076-3417/10/20/7293/s1>, Supplementary Material S1: “specifications on multi-GAS infrared (IR) and electrochemical sensors and gas standards used for instrument calibration” is available for consultation.

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