



Magma source, pre-eruptive dynamics and timescales of major explosions at Stromboli volcano (Italy)

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Abstract

At Stromboli volcano (Italy) the regular, persistent activity is occasionally (2–4 events/year) interrupted by sudden, short-lived and more energetic major explosions, being intermediate in size between regular Strombolian activity and even more powerful paroxysmal explosions. Despite being frequent and hazardous, the magma source depth, the triggering mechanisms and timescales of such major explosions are still poorly understood. Here, we focus on three major explosions which occurred on 3 May, 8 November and 24 November 2009. We present a dataset of major element composition and dissolved volatile contents in olivine-hosted melt inclusions, embayments and glassy groundmass. We combine them with Fe–Mg diffusion profiles in olivine phenocrysts, and with volcanic gas plume and ground tilt measurements. Olivine phenocrysts display Fo_{69-72} compositions, with reverse zoning (up to Fo_{83}) in the 24 November and to a minor extent in the 8 November eruption. Glassy groundmass of the November events ranges from the more evolved towards more primitive compositions, whereas the 3 May 2009 glassy groundmass has evolved compositions. Dissolved H_2O and CO_2 in glassy and bubble-free melt inclusions are low, with CO_2 below the detection limit and H_2O up to 2.3 wt.%. Volcanic plume measurements indicate the occurrence of these events in phases of high CO_2 fluxes, with substantial geochemical changes preceding the onset of the 24 November event. Based on these results, we propose that the 3 May major explosion — the smallest in intensity/magnitude — was likely driven by the accumulation of gas bubbles into the shallower (< 1 km below sea level, b.s.l.) HP reservoir. In contrast, we find the November major explosions to be triggered by the injection of variable amounts of deeper-stored LP magma and gas into the HP reservoir, over hours to weeks before.

Keywords Explosive basaltic eruptions · Stromboli · Major explosions · Melt inclusions · Olivine · Volcanic plume chemistry

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Introduction

Basaltic, open-vent volcanoes are prominent sources of volcanic gases into the atmosphere (Carn et al. 2016; Edmonds et al. 2022), and produce a wide spectrum of explosive eruptions, varying in scale and style from low-intensity Strombolian eruptions to paroxysmal events of Vulcanian to Plinian magnitude (e.g., Houghton and Gonnermann 2008; Allard 2010; Vergnolle and Métrich 2021; Barth et al. 2024; Vergnolle 2024).

Stromboli, located at the north-easternmost edge of the Aeolian archipelago (Southern Italy) (Fig. 1a), is one of these open-vent volcanoes (Aiuppa et al. 2021; Métrich et al. 2021; Ripepe et al. 2021), having been characterized by open-vent conditions at least since the seventh century CE (Rosi et al. 2013).

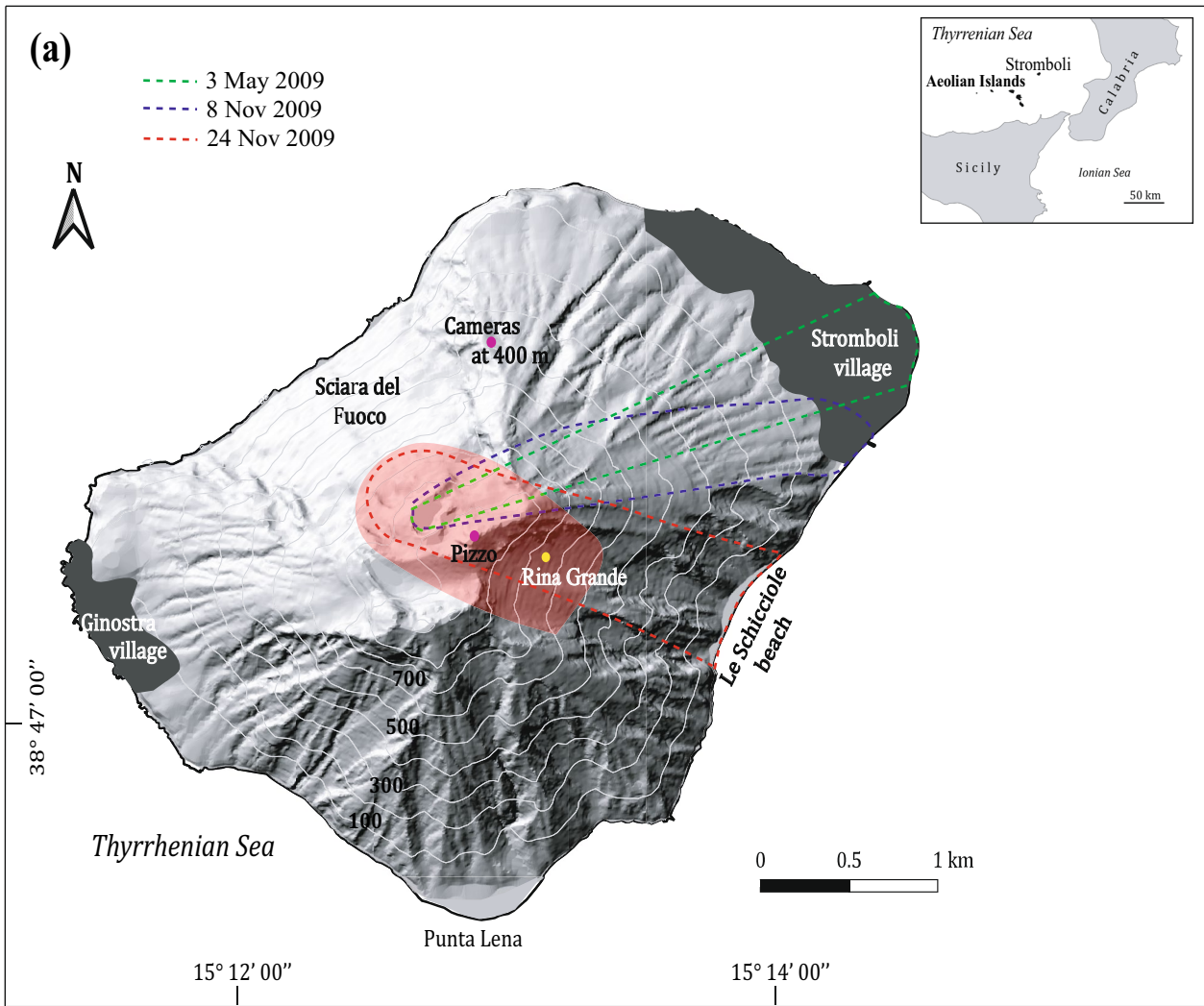


Fig. 1 a Shaded relief map of Stromboli Island showing the tephra dispersal of the 3 May, 8 November and 24 November 2009 major explosions (dashed green, blue and red lines, respectively), the location of video-cameras and of the main localities. Red area indicates the area affected by spatters and bombs during the 24 November major explosion. Data of products dispersion are taken from Andronico (2009) for the 3 May 2009 and from Andronico and Pistolesi (2010) for the 8 and 24 November 2009 major explosions. **b** 3 May, **c** 8 November and **d** 24 November 2009 eruptive products. Note the mingled bomb showing the black scoriaceous HP and the LP pumiceous components

Its persistent explosive dynamics is characterized by rhythmic, mild to moderate Strombolian explosions, known as *ordinary activity*. These explosions repeat every 10–20 min on average, last a few seconds, and eject pyroclastic material up to heights of 100–200 m (Ripepe et al. 2008; Taddeucci et al. 2012; Kelfoun et al. 2020). Tephra volumes are typically $\sim 1\text{--}10\text{ m}^3$, with mass discharge rates of $10^2\text{--}10^3\text{ kg/s}$ (Rosi et al. 2013 and references therein). Such regular explosive activity is preceded by a $\sim 150\text{ s}$ long, low-amplitude ($\sim 0.1\text{ }\mu\text{rad}$) inflation of the ground and is followed by a rapid deflation ($\sim 30\text{ s}$) (Genco & Ripepe 2010; Ripepe et al. 2021).

This steady activity is occasionally interrupted by sudden, more violent and short-lived (tens of seconds to few minutes) explosions often involving multiple vents simultaneously and ranging in intensity and magnitude from *major explosions* to *paroxysms* (Barberi et al. 1993; Bertagnini et al. 2011). The last paroxysms occurred in 2003, 2007, 2019 and 2024 (Métrich et al. 2005, 2010, 2021; Andronico et al. 2021; Giordano and De Astis 2021) — they are highly energetic (mass discharge rate $> 10^6\text{ kg/s}$; Rosi et al. 2006, 2013; Pistolesi et al. 2011; Pioli et al. 2014) explosions and produce eruptive plumes rising to 3–8 km above the crater terrace (Pioli et al. 2014) or up to 10 km in the larger-scale events (e.g., 22 May 1919 or 11 September 1930; Ponte 1919; Rittman 1931). Paroxysms are associated with the ejection of bombs and meter-sized ballistic blocks at distances up to 2.5 km from the vents. These more violent events are preceded by longer (up to 600 s) and larger ($\sim 10\text{ }\mu\text{rad}$) ground deformation compared to the ordinary activity.

Extensive geochemical and petrological research has been conducted on pyroclastic materials erupted at Stromboli, finding diverse compositional signatures for ordinary activity and paroxysms.

Ordinary activity is fed by a crystal-rich ($\sim 50\text{ vol.}\%$) and partially degassed ($\text{H}_2\text{O} < 0.6\text{ wt.}\%$, $\text{CO}_2 < 100\text{ ppm}$, $\text{S} < 1300\text{ ppm}$, $\text{Cl} < 2700\text{ ppm}$) basaltic-shoshonitic magma ($\text{K}_2\text{O} = 1.9\text{--}2.5\text{ wt.}\%$; $\text{SiO}_2 = 48.5\text{--}51.5\text{ wt.}\%$), emitted as a high-porphyrific (HP) black scoria. The mineral assemblage consists of phenocrysts of plagioclase ($\text{An}_{60\text{--}90}$), clinopyroxene ($\text{Mg}\#_{0.70\text{--}0.90}$, $\text{Fs}_{5\text{--}17}$), olivine ($\text{Fo}_{70\text{--}74}$), all set in a shoshonitic glassy groundmass ($\text{K}_2\text{O} > 3.8\text{ wt.}\%$; Bertagnini et al.

2008; Francalanci et al. 2004; Métrich et al. 2001, 2010; Landi et al. 2004, 2008, 2022).

In contrast, paroxysms are fed by a volatile-rich basaltic magma ($\text{H}_2\text{O} = 1.8\text{--}3.4\text{ wt.}\%$; $\text{CO}_2 = 890\text{--}1890\text{ ppm}$; $\text{S} = 1660\text{--}2250\text{ ppm}$; $\text{Cl} = 1660\text{--}2030\text{ ppm}$) straddling the high-K (HK) shoshonitic field. This is erupted as crystal-poor ($< 10\text{ vol.}\%$) and low-porphyrific (LP) golden pumices. Mineralogy consists of clinopyroxene ($\text{Mg}\#_{0.84\text{--}0.90}$, $\text{Fs}_{5\text{--}8}$) and olivine with predominant composition $\text{Fo}_{85\text{--}86}$ in small-scale paroxysms (e.g., 15 March 2007, 3 July 2019) and $\text{Fo}_{88\text{--}91}$ in the larger-scale paroxysms (e.g., 11 September 1930), set in a shoshonitic-basaltic, vesicle-rich glassy groundmass ($\text{K}_2\text{O} < 2.4\text{ wt.}\%$; Bertagnini et al. 2003, 2008; Francalanci et al. 2004; Landi et al. 2004, 2009, 2022; Métrich et al. 2001, 2005, 2010, 2021).

Based on these evidence, previous studies have inferred the existence of a complex, vertically extended plumbing system.

The ordinary activity is caused by the surface bursting of gas slugs, formed by the coalescence of smaller bubbles within the upper conduit ($< 3\text{ km b.s.l.}$; Burton et al. 2007) resident HP magma.

Paroxysms instead have been recognized to have a deeper source, and are triggered by the interplay of several processes including (i) the intrusion of hot and CO_2 -rich bubbly magma causing pressurization of the deep volcano's plumbing system (7–11 km b.s.l.) (Métrich et al. 2001, 2005, 2021; Bertagnini et al. 2003, 2008; Francalanci et al. 2004; Pichavant et al. 2011), (ii) collapse and fast ascent of a deeply accumulated CO_2 -rich foam, previously stored at the roof of the LP reservoir (Allard 2010; Aiuppa et al. 2011, 2021) or at some rheological discontinuity (Caricchi et al. 2024).

Based on eruptive parameters, paroxysms can be also separated in two main categories (Large-scale –LSP– and Small-scale –SSP– paroxysms), which in turn reflect slight variability of magmatic source region and compositional characteristics (Bertagnini et al. 2008, 2011; Métrich et al. 2005, 2010, 2021; Pistolesi et al. 2011; Rosi et al. 2013).

Constraints on the timescales of paroxysms are given by Fe–Mg diffusion profiles in olivine crystals and suggest magma ascent times of 2–10 days from 6 to 9 km to the surface, and long (months) incubation times for destabilization of the LP magma reservoir (Métrich et al. 2021). These timescales are in agreement with timescales derived from plume gas measurements (Aiuppa et al. 2021).

In contrast to ordinary activity and paroxysms whose magma sources, pre-eruptive dynamics and timescales are relatively well constrained, much less is known on the triggering mechanisms of major explosions.

Here, we present a petrological and geochemical study of three major explosions of variable intensity occurred on 3 May, 8 November and 24 November 2009, with the aim of

reconstructing their pre-eruptive dynamics and their timescales from the deep storage level to ascent to surface. To this aim (i) we retrieve the pre-eruptive storage conditions by measuring dissolved volatiles contents (H_2O , CO_2 , S, Cl) in olivine hosted melt inclusions, accompanied by analysis of major and minor element of their host olivine crystals, and (ii) we constrain the timescales of pre-eruptive processes by modelling Fe–Mg compositional profiles in zoned olivine crystals.

Source and dynamics of major explosions: state of the art

Major explosions are sporadic (2 events/year; Barberi et al. 1993; Bertagnini et al. 2008; Bevilacqua et al. 2020) eruptions of intermediate size (Pioli et al. 2014). They produce plumes a few hundred metres in height (Bertagnini et al. 2003; Andronico et al. 2008) forming discontinuous blankets of tephra deposits sporadically reaching the coastline, and a shower of bombs and meter-sized lithic blocks covering the upper slopes of the volcano, typically to maximum distance of ~ 1.5 km from the vents.

The ejected tephra volumes are typically in the 10^2 – 10^3 m^3 range, with mass discharge rates of 10^4 – 10^5 kg/s (Rosi et al. 2013; Pioli et al. 2014). These mid-intensity events are preceded by smaller (~ 1 μrad) ground deformation compared to paroxysms, starting ~ 300 s before the onset of the eruption. Nevertheless, they are characterized by similar deformation trends (Ripepe et al. 2021).

Major explosions are characterized by a heterogeneity of their eruptive products. The lowest-intensity explosions produce exclusively HP scoria (e.g., 8 September 1998; Bertagnini et al. 1999), as well as the ordinary activity. The higher intensity events (3 May, 8 November, 24 November 2009; La Felice and Landi 2011) are instead complex mixtures of both HP scoria and LP pumices, also showing intermediate glassy groundmass composition in between.

In addition to the heterogeneity of their eruptive products, interpretation is additionally complicated by the under-sampling of these events, which deposits are poorly dispersed and usually not long preserved.

A relatively deeper source origin than ordinary activity is supported by the gas observations (Aiuppa et al. 2021) that recorded gas plume CO_2/SO_2 ratios above background in six major explosions occurred between 2018 and 2020, including the 19 July 2020 event. Geophysically, this event exhibited a ground tilt ~ 4 times higher than the average for major explosions (3.5 μrad vs. 0.8 μrad), thus ranking at the boundary between major explosions and small-scale paroxysms (Ripepe et al. 2021).

Voloschina et al. (2023) used H_2O and CO_2 compositions measured in olivine-hosted melt inclusions to constrain

the magma source of this peculiar major explosion to be at ~ 9.5 km b.s.l. depth with the activation of a shallower (5–6 km b.s.l.) ponding zone. Based on Fe–Mg diffusion profiles in olivine crystals, they suggested shorter incubation (20–25 days before the onset) with longer ascent times (5–10 days) than those associated to paroxysms. However, it is currently unknown if similar triggering/incubation depths and timescales also apply to other, less energetic, major explosions.

The 2009 major explosions at Stromboli

Three major explosions hit Stromboli on 3 May, 8 November and 24 November 2009, with the latter event being the most energetic since the 2007 paroxysm for erupted volume, pyroclastic tephra dispersal and recorded amplitude of syn-eruptive seismic and infrasonic signals as recorded by the various monitoring networks deployed on the island (Andronico and Pistolesi 2010; Aiuppa et al. 2011).

The 3 May 2009 event was preceded by deformation of the summit crater area since 20 March 2009 as detected by ground-based interferometric synthetic aperture radar (Nolesini et al. 2013). This ground deformation episode reached a maximum of 0.35 mm/h on 27 March 2009 and remained high for the following month until 3 May 2009, when it was 0.25 mm/h. INGV (Istituto Nazionale di Geofisica e Vulcanologia) reported a first explosion at 14:58:08 GMT (Cristaldi 2009a) from the central sector of the crater terrace, producing the emission of coarse pyroclasts that reached a height of 200 m above the crater rim, and then fell mostly within the crater terrace but also outside its NW sector, and was followed by a terminal ash emission. As observed from visible and thermal cameras, a second larger explosion occurred at 14:58:28 GMT from the southern vent, radially ejecting metric-sized dark spatter bombs over the whole field of view of a camera installed on the summit area at Pizzo (Fig. 1a). The eruptive plume generated a shower of ash and lapilli in ENE direction, eventually reaching the village of Stromboli and forming a discontinuous tephra deposit that was partially washed away by a violent storm the day after (Andronico 2009). Lava fountaining and ash emission were observed to follow the main event from other vents. An intense spattering activity started at 15:00 GMT and produced a small intracrateric lava flow (Andronico 2009). The monitoring network of LGS (Laboratorio di Geofisica Sperimentale – University of Florence, Italy n.d.) recorded an associated seismic, very-long-period (VLP) amplitude of 7 μm , a ground deformation of ~ 0.8 μrad and an infrasonic pressure of ~ 5 bar, thus classifying the 3 May 2009 event as a major explosion.

After the 3 May event, a new increase in the deformation rate at the summit crater area started on 2 November

2009 (0.56 mm/h) as recorded by the ground-based interferometric synthetic aperture radar and reached 0.6 mm/h on 8 November 2009 (Nolesini et al. 2013) when, at 12:29:31 GMT, a mild explosion (column height, ~ 50 m) occurred at the central crater (Cristaldi 2009b). This produced the fallout of coarse, incandescent materials and was followed by two main explosions of lapilli and ash, that lasted about 38 s. Tephra was dispersed ENE within a distance of 2.5 km from the vents, covering a dispersal area on the ground of around 10,000 m² (Andronico and Pistolesi 2010). Intense lava fountaining from the southern sector started at 12:30:25 and lasted 50 s, followed by 20 min of intense continuous spattering from the central sector. The activity was accompanied for 5 min by the effusion of an intracrateric lava and was classified as a major explosion. The LGS monitoring network recorded a seismic velocity and displacement of 8×10^{-4} m/s and 3×10^{-5} m, respectively, 6 bar of infrasonic pressure and a ground deformation of 0.57 μ rad.

After 8 November, activity returned to background levels but on 24 November 2009, another major explosion occurred at 11:20:48 GMT, consisting of simultaneous bursts from two active vents located in the southern sector of the crater associated with a recorded ground deformation of 0.61 μ rad recorded by the LGS network. A ~ 150 m eruptive plume generated ash to lapilli fallout, also ejecting proximal bombs and lava fragments from a central, previously inactive vent. A second, more intense explosion followed after 15 s that radially ejected a larger volume of juvenile bombs and lithic clasts, affecting the whole summit area at least 300 m from the vent. The eruptive plume emplaced tephra fallout over the summit area and in particular in the Pizzo sopra la Fossa area and towards E reaching the shoreline at the Schicciole beach, 1.6 km SE from eruptive vent (Fig. 1a). Several fires were ignited by tephra fallout in the eastern sector of the island below Rina Grande. The second explosion produced a 50 m wide crater in the central sector of the crater terrace, where a following intense degassing and spattering were observed. The total sequence lasted 50 s and produced a larger dispersal and volume of tephra compared to the other two major explosions, covering an area of 15,000 m² (Andronico and Pistolesi 2010).

Sample preparation and analytical methods

Criteria of sample selection and description

Samples were collected by D. Andronico and M. Pistolesi few days after each explosive event (see Andronico and Pistolesi 2010 for details). Mingled pumices were crushed and sieved in the 1–0.5 mm and 0.5–0.25 mm granulometric fractions. Olivine crystals were hand-picked under a binocular stereomicroscope and were embedded in orthodontic

resin. A total of 57 melt inclusions (MIs) and 3 embayments were selected and doubly polished in order to expose both sides, except some exceedingly thin (< 10 μ m) melt inclusions that were exposed only on one side.

Melt inclusions were found to be more abundant in olivine relative to pyroxene or plagioclase crystals; we hence selected and analysed exclusively those hosted in olivine crystals. Olivine-hosted melt inclusions have sizes ranging from 20 to 180 μ m in diameter, and show variable morphologies: well-rounded, ellipsoidal, faceted (negative host crystal shapes), squared and irregularly shaped (Fig. 2a–c; Fig. S1 in Supplementary Figures).

After their entrapment, melt inclusions may be affected by modifications in response to changing T-P-X-fO₂ magma conditions, including crystallization of the host mineral at the host-inclusion interface and exsolution of volatiles into a bubble (shrinkage bubble), that can significantly deplete CO₂ in the melt due to its low solubility (Anderson and Brown 1993; Dixon and Stolper 1995; Métrich and Wallace 2008; Wallace et al. 2015a,b; Rose-Koga et al. 2021). The majority (48/57) of melt inclusions analysed in this study is entirely glassy, whereas a minority contain mineral daughters (4/57) or bubbles (3/57); relatively to these latter, bubble volumes are all in the usual range of shrinkage bubbles (0.2 to 5 vol% of the melt volume; Lowenstern 1995; Steele-MacInnis et al. 2017), excluding the heterogeneous trapping of a mixture consisting of gas and melt mixture.

A description of analysed melt inclusions and host olivine crystals, from each eruptive event, is reported in Table S1 (Supplementary Material 1) detailing location of melt inclusions in olivine crystals (core/rim), size, shape, eventual presence and size of bubbles and/or daughter minerals. Embayments and glassy groundmass wetting the olivine phenocrysts were also analysed.

Fourier-transform infrared spectroscopy

Fourier-transform infrared spectroscopy (FTIR) was conducted at the Dipartimento di Scienze della Terra of the University of Pisa (Italy) using a Nicolet iN10 FTIR Microscope equipped with a global infra-red source and a Ge-coated KBr beamsplitter, in order to analyse dissolved volatiles (H₂O, CO₂) in melt inclusions, embayments and glassy groundmass. For spot analyses, a liquid-nitrogen cooled MCT detector was used, and the aperture varied from 11 to 33 μ m square, depending on inclusion size. In larger melt inclusions, several spectra were acquired using different sizes to verify the consistency of aperture size, in agreement with the recommendations of Rose-Koga et al. (2021). Spectra were acquired in the 400–6000 cm⁻¹ absorption range, collecting 512 scans at a resolution of 8 cm⁻¹ for glasses and 256 scans for olivine crystals and background. The beam path within the microscope

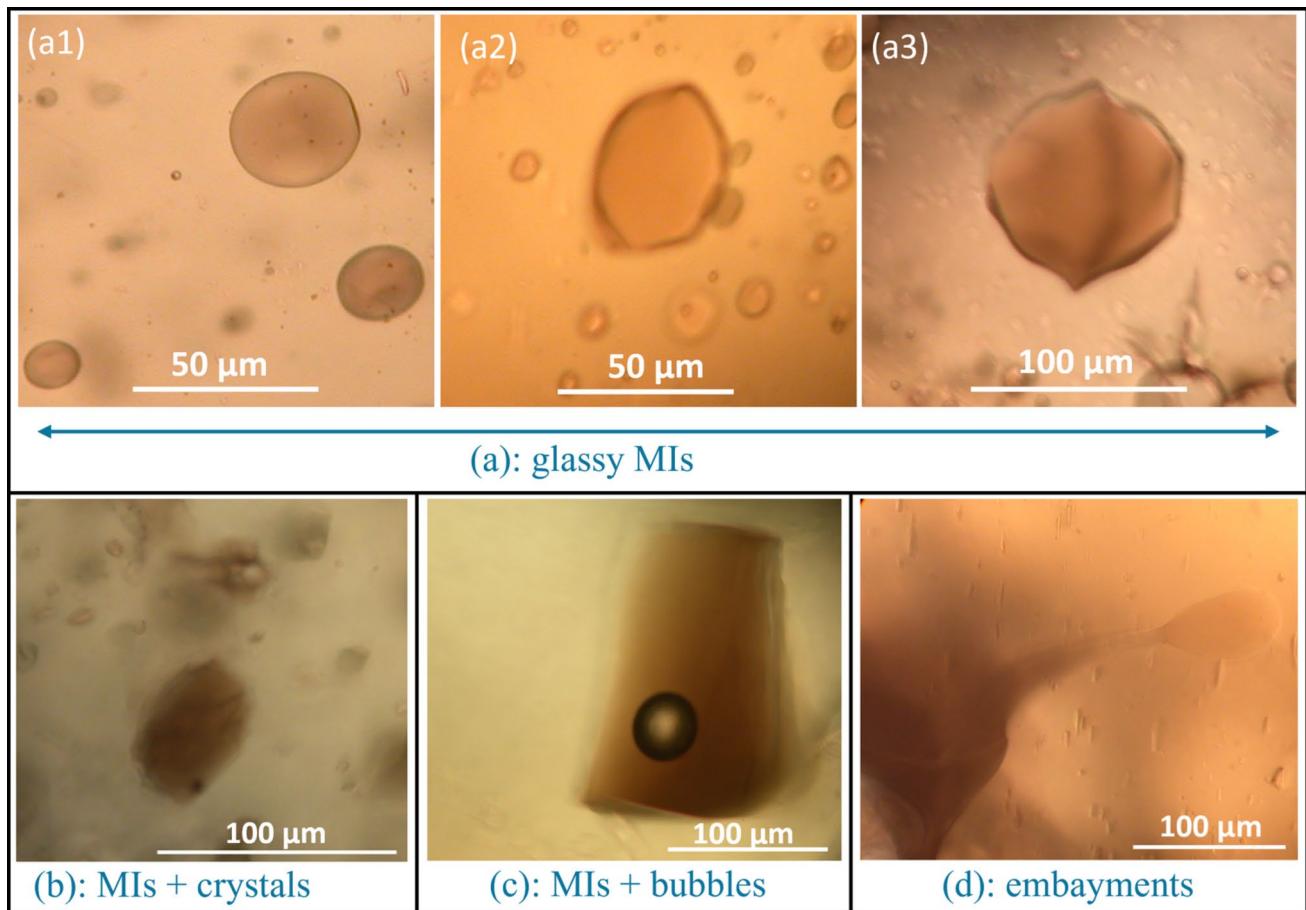


Fig. 2 Transmitted light microphotographs of olivine-hosted melt inclusions with various morphologies: (a): glassy melt inclusions; (a1): well-rounded ($a=b$ axis), glassy melt inclusions; (a2) faceted,

large melt inclusion; (a3) squared melt inclusion; (b): melt inclusion with mineral daughters; (c): melt inclusion with a 40 μm shrinkage bubble; (d) melt embayment

was continuously purged with a continuous flux of dry, CO_2 -free compressed air with a -73 $^\circ\text{C}$ dewpoint. The software Omnic Picta was used to control the microscope's imaging and FTIR capabilities.

H_2O and CO_2 concentrations were calculated according to the Beer-Lambert law:

$$C = \frac{100 \times \text{Abs} \times M}{\epsilon \times \rho \times t}$$

where Abs is the absorbance, M the molar mass (g/mol), ϵ the extinction coefficient (L/mol per cm), ρ the glass density, and t the thickness (cm). Water absorption was calculated by measuring the total H_2O absorption band centred at a wavenumber of 3535 cm^{-1} using a straight baseline correction, while the CO_3^{2-} peak doublet was collected at 1520 cm^{-1} and 1430 cm^{-1} . CO_2 peak deconvolution was carried out through Peak Fit software after the subtraction of a degassed glass spectrum obtained from measurements of the glassy groundmass. The partially exposed melt inclusions and embayments were corrected for olivine crystal

spectra, by measuring olivine absorption between 1600 and 2000 cm^{-1} in both the glass and in the host mineral spectra (Von Aulock et al. 2014). Density used is an averaged value (2.69 ± 0.02 g/cm^3), determined on Stromboli hydrous glasses (Métrich et al. 2001). Extinction coefficients (ϵ) were: $\epsilon^{3535} = 64.3$ L/mol per cm (Métrich et al. 2001) and $\epsilon^{1520} = 362$ L/mol per cm (Voloschina et al. 2023). Olivine thickness (t) was previously measured with a calibrated microscope stage at magnifications of $10\times$ and $50\times$. Multiple measurements were taken at different spots along the crystals and an average thickness was then calculated for individual crystals. Analytical uncertainties in H_2O and CO_2 concentrations were calculated based on propagation of error from wafer thickness, determining a maximum error of 13%. Given that H_2O and CO_2 solubilities are strongly dependent on pressure, under the assumption that melt inclusions were vapor-saturated at the time of entrapment, the use of a H_2O - CO_2 solubility model (i.e., Newman and Lowenstern 2002; Iacono-Marziano et al. 2012; Ghiorso and Gualda 2015; Shishkina et al. 2014) allowed calculating their

saturation pressures (P_{sat}) and, thus, to retrieve the magmatic storage depth (e.g., Métrich and Wallace 2008; Wallace et al. 2021). Calculations were done in VESICAL 1.0.1 (Iacovino et al. 2021), using a JupyterLab computing environment hosted on the ENKI server (<http://enki-portal.org/>).

Electron microprobe analyses

Samples previously analysed by FTIR and additional olivine crystals were embedded in epoxy resin and then carbon coated for microanalysis. Major element, S and Cl composition of glasses (melt inclusions, embayments and glassy groundmass) and olivine crystals were obtained by electron microprobe analysis (EPMA) at the joint laboratory (LaMA) of the DST and CNR-IGG of Firenze (Italy) using a JEOL Superprobe JXA-8230 equipped with 5 wavelength dispersion spectrometers (WDS). Glasses and olivine crystals from 3 May 2009 were analysed at the Unitech COSPECT at the University of Milan, Italy, using a JEOL 8200 Super Probe. Analyses on olivine crystals were performed using an accelerating voltage of 15 kV, a beam current of 20 nA and a beam diameter of 3 μm . Glass analyses were performed at 15 kV, 10 nA, using a defocused beam diameter of 5 to 10 μm . ZAF correction for matrix effects was applied. Different counting times were set for major and minor elements: 15 s on peak and 7 s on background for major elements, 30–40 s on peak and 15–20 s for minor elements, excluding Na. To reduce alkali loss, Na was measured first, for 10 s on peak and 5 s on background. The relative elemental average errors were obtained from counting statistics and were found to be $\leq 2\%$ for Mg, $\leq 2\%$ for Fe, 1% for Ca, Al and K, $\leq 3\%$ for Na, 5% for Cl (9% for the 3 May products) and 27% for S (the latter increasing in analyses of the glassy groundmass). The analytical standards used for calibration were basaltic glasses from SE Indian ocean (NMNH 113716–1, NMNH 111240–52). Analytical errors and accuracy of electron microprobe analyses are reported in Table S11–15 (Supplementary Material 3).

In the larger melt inclusions, several spot analyses were collected and then averaged. A total of 63 spot analyses were collected on the glassy groundmass, but specifically on the 3 May 2009 clasts, the low amount of the LP component did not allow a detailed analysis of its glassy groundmass and crystals. Olivine composition was analysed in multiple spots close to MIs and in the rims of host crystals, for a total of 240 spot analyses collected. Due to the double polishing procedure often leaving only olivine fragments, additional olivine phenocrysts were selected, oriented and mounted in epoxy resin for collecting Fe–Mg diffusion profiles, obtaining a total of 15 concentration profiles. Diffusion profiles were positioned following the suggestions of Shea et al. (2015).

Post-entrapment crystallization and melting

Post-entrapment crystallization (PEC) and melting (PEM) were calculated by using the Melt Inclusion Modification Correction (MIMiC) program (Rasmussen et al. 2020), which calculates the olivine-melt equilibrium Fe–Mg distribution coefficient ($K_d^{\text{Fe-Mg}}_{\text{ol-melt}}$) using the model of Toplis (2005), then adding incrementally olivine composition to the melt inclusion until equilibrium K_d is reached.

Post-entrapment crystallization (PEC) of olivine mainly affects MgO, FeO and SiO_2 , which are compatible in olivine, whereas incompatible elements (e.g. volatile elements) could suffer a potential increase only if the amount of PEC is $> 5\%$ (Rose-Koga et al. 2021). The amount of PEC required for the analysed MIs is $< 5.5 \text{ wt.}\%$ (Table S2, Supplementary Material 1; Fig. S10 in Supplementary Figures). Melt inclusions may also undergo post-entrapment melting due to overheating (Sobolev and Kostyuk 1975; Rasmussen et al. 2020), but this effect is limited in our case (1.5 wt.% on average; Table S2, Supplementary Material 1). Considering that both PEC and PEM values are negligible and that ratios such as $\text{CaO}/\text{Al}_2\text{O}_3$ and $\text{K}_2\text{O}/\text{Na}_2\text{O}$, as well as H_2O , CO_2 and S would be diluted to a similar degree, in the following not-re-calculated compositions are considered.

Fe–Mg timescales

A total of 10 out of 15 Fe–Mg concentration profiles in olivine phenocrysts (6 from the 24 November and 4 from the 8 November 2009) were modelled by using the Diffusion Process Analysis (DIPRA) software (Girona and Costa 2013), in order to calculate timescales of pre-eruptive magmatic processes.

The orientation of olivine crystals was carried out by visual inspection of the crystal shape under an optical microscope.

Pre-eruptive conditions were fixed at 1085 $^{\circ}\text{C}$ (± 37 $^{\circ}\text{C}$) and 76 MPa. The chosen temperature represents an average of values calculated using the geothermometer of Putirka et al. (2007) on 14 melt inclusions from this study. Although it is lower than the 1150 $^{\circ}\text{C}$ temperature used for Fe–Mg diffusion calculation for the 2020 major explosion and for paroxysms (Voloschina et al. 2023; Métrich et al. 2021), it still falls within the lower temperature range (1050–1175 $^{\circ}\text{C}$) derived from experimental studies conducted on a LP basalt pumice of Stromboli (Di Carlo et al. 2006). The pressure value of 76 MPa was chosen in order to simulate the shallow magmatic reservoir of Stromboli, in accordance with the lower pressure bound (50–100 MPa) reported by Di Carlo et al. (2006). The redox conditions were fixed at NNO + 1 (NNO refers to the Ni–NiO oxygen buffer), in line with previous studies (e.g., Aiuppa et al. 2010; Métrich et al. 2021; Voloschina et al. 2023) and representative of the oxidizing

conditions of the Stromboli magmas. Natural concentration profiles were modelled considering non-homogeneous and step initial conditions. Input parameters used in DIPRA are reported in Table S10 (Supplementary Material 2).

Volcanic gas

Volcanic gas data were obtained from daily survey of the plume flux of Stromboli volcano carried out during the 2006–2010 years. Specifically, the CO_2 plume flux was determined by simultaneous measurements of the CO_2/SO_2 plume ratio by three fully automated Multi-GAS instruments and SO_2 mass flux was determined by a remotely controlled network consisting of four UV scanning DOAS spectrometers (see Aiuppa et al. 2011 for further details and for location of the MultiGAS and UV scanner stations).

Results

Major element data

Groundmass composition

Glassy groundmass from the 24 November 2009 samples ranges from shoshonitic ($\text{K}_2\text{O} = 2.08\text{--}2.48$ wt.%; $\text{SiO}_2 = 48.71\text{--}50.80$ wt.%) to shoshonitic-basaltic ($\text{K}_2\text{O} = 3.59\text{--}3.94$ wt.%; $\text{SiO}_2 = 49.83\text{--}51.83$ wt.%) (Fig. S2 in Supplementary Figures), in agreement with previous results reported in La Felice and Landi (2011). These compositions cluster both in the fields of LP ($\text{K}_2\text{O} < 2.4$ wt.%) and HP ($\text{K}_2\text{O} > 3.8$ wt.%) magmas, respectively (Landi et al. 2022). The high variability in K_2O contents likely suggests mixing between magma batches having slightly variable melt and mineral compositions (Voloschina et al. 2023). This is further observable in Fig. 3 and 4 where major element ratios exhibit a wide variability ($\text{K}_2\text{O}/\text{Na}_2\text{O} = 0.77\text{--}1.20$; $\text{CaO}/\text{Al}_2\text{O}_3 = 0.48\text{--}0.67$), spanning from a more evolved composition ($\text{K}_2\text{O}/\text{Na}_2\text{O} > 1$; $\text{CaO}/\text{Al}_2\text{O}_3 < 0.5$) to the compositional fields of the April 2003 and March 2007 small-scale paroxysms (SSP; Métrich et al. 2005, 2010) and historic large-scale paroxysms (LSP; Métrich et al. 2010).

For comparison, the glassy groundmass of the 19 July 2020 major explosion (Voloschina et al. 2023) shows a similar compositional trend. The 24 November 2009 glassy groundmass is Fe-richer ($\text{FeO} = 7.77\text{--}9.77$ wt.%; $\text{MgO} = 3.58\text{--}5.83$ wt.%; $\text{FeO}/\text{MgO} = 1.44\text{--}2.70$) than the groundmass of 19 July 2020 ($\text{FeO} = 7.32\text{--}8.97$ wt.%; $\text{MgO} = 4.21\text{--}7.68$ wt.%; Voloschina et al. 2023). The glassy groundmass of the 8 November 2009 eruption shows intermediate compositions, from shoshonitic basalt toward the shoshonitic basaltic field straddling with the HK basalt field ($\text{K}_2\text{O} = 2.23\text{--}4.06$ wt.%; $\text{MgO} = 3.49\text{--}7.45$ wt.%;

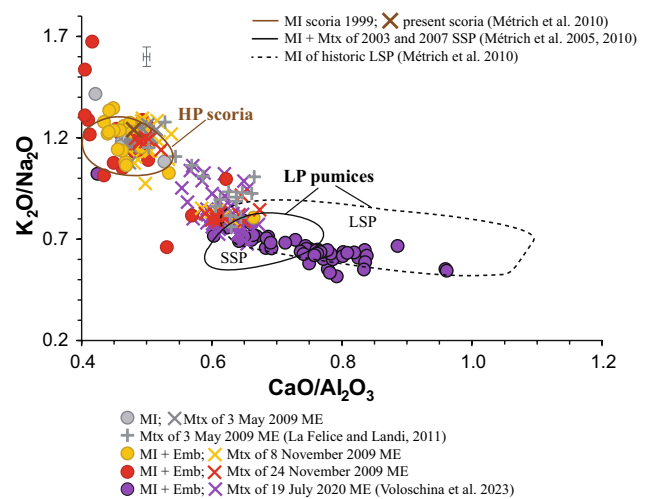


Fig. 3 $\text{K}_2\text{O}/\text{Na}_2\text{O}$ vs $\text{CaO}/\text{Al}_2\text{O}_3$ ratios of melt inclusions (MIs), embayments (Emb) and glassy groundmass (Mtx) from different Stromboli eruptions: 1999 scoria (ordinary activity, Métrich et al. 2010); 3 May 2009 (this study; La Felice and Landi 2011), 8 November 2009 (this study) and 24 November 2009 (this study) major explosions (ME); 19 July 2020 major explosion (Voloschina et al. 2023); 2003 and 2007 small-scale paroxysms (SSP; Métrich et al. 2005, 2010); historic large-scale paroxysms (LSP; Métrich et al. 2010). Filled dots represent melt inclusions and/or embayments, cross symbols represent glassy groundmass. Average errors ($\text{K}_2\text{O}/\text{Na}_2\text{O} = 3\%$ and $\text{CaO}/\text{Al}_2\text{O}_3 = 1\%$) are indicated as error bars. HP, high porphyritic; LP, low porphyritic

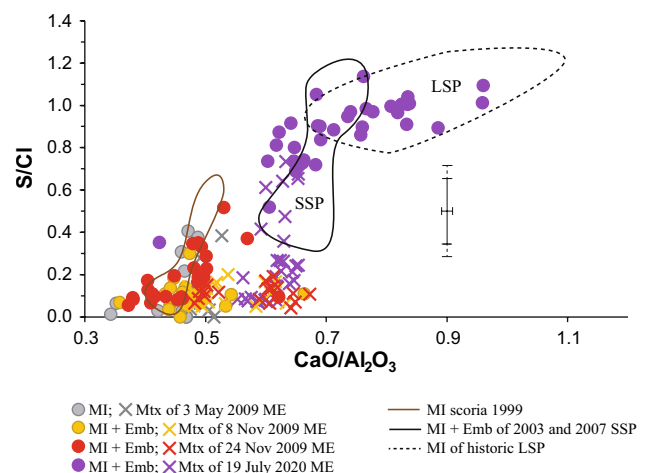


Fig. 4 S/Cl vs $\text{CaO}/\text{Al}_2\text{O}_3$ ratios of melt inclusions, embayments and glassy groundmass from different eruptions at Stromboli: 1999 scoria (ordinary activity, Métrich et al. 2010); 3 May 2009, 8 November and 24 November 2009 major explosions (ME; this study); 19 July 2020 major explosion (Voloschina et al. 2023); 2003 and 2007 small-scale paroxysms (SSP; Métrich et al. 2005; 2010); historic large-scale paroxysms (LSP; Métrich et al. 2010). Filled dots represent melt inclusions (MI) and/or embayments (Emb), crosses represent glassy groundmass (Mtx). Average errors ($\text{S}/\text{Cl} = 31\%$ for MI/Emb and 43% for Mtx; $\text{CaO}/\text{Al}_2\text{O}_3 = 1\%$) are indicated as error bars: solid bar indicates MI and Emb errors, dashed bar indicates Mtx error

$\text{SiO}_2 = 48.06\text{--}52.09$ wt.%) (Fig. S2 in Supplementary Figures). Glassy groundmass of the 3 May 2009 eruption is shoshonitic-basaltic in composition ($\text{K}_2\text{O} = 3.78\text{--}4.04$ wt.%; $\text{MgO} = 3.39\text{--}3.88$ wt.%, $\text{SiO}_2 = 49.89\text{--}53.64$ wt.%) and plots in the HP compositional field (Fig. 3, $\text{K}_2\text{O}/\text{Na}_2\text{O} = 1.08\text{--}1.27$; $\text{CaO}/\text{Al}_2\text{O}_3 = 0.47\text{--}0.51$).

Melt inclusions and embayments

Major element ratios in melt inclusions and embayments of the 24 November 2009 ($\text{K}_2\text{O}/\text{Na}_2\text{O} = 0.66\text{--}1.67$; $\text{CaO}/\text{Al}_2\text{O}_3 = 0.37\text{--}0.62$) mostly overlap the HP magma compositions, except few MIs that have intermediate compositions close to the LP compositional field. Melt inclusions and embayments are even Fe-richer ($\text{FeO} = 9.05\text{--}12.73$ wt.%; $\text{MgO} = 2.62\text{--}3.99$ wt.%; $\text{FeO}/\text{MgO} = 2.51\text{--}4.52$) than associated groundmass and melt inclusions from July 2020 ($\text{FeO} = 5.93\text{--}9.17$ wt.%, $\text{MgO} = 1.94\text{--}5.52$; Voloschina et al. 2023), 5 April 2003 ($\text{FeO} = 7.43\text{--}9.09$ wt.%; $\text{MgO} = 4.99\text{--}6.78$ wt.%; Métrich et al. 2005), and 15 March 2007 ($\text{FeO} = 7.49\text{--}8.43$ wt.%, $\text{MgO} = 4.85\text{--}5.12$ wt.%; Métrich et al. 2010). According to Bertagnini et al. (2003), Fe-rich melt inclusions could derive from the entrainment and dissolution of pre-existing Fe-richer olivine crystals into melts of different differentiation degree, with successive crystallization and possible post-entrapment re-equilibration. Generally, composition of glasses is comparable but more evolved than whole-rock compositions of the same eruption ($\text{MgO} = 6.07\text{--}6.02$ wt.%, $\text{CaO} = 11.21\text{--}11.46$; La Felice and Landi 2011).

Major elements ratios in melt inclusions of the 8 November 2009 ($\text{K}_2\text{O}/\text{Na}_2\text{O} = 0.79\text{--}1.30$, $\text{CaO}/\text{Al}_2\text{O}_3 = 0.47\text{--}0.63$) cover the entire range from the HP toward the LP compositional fields (Fig. 3). Compositions cluster in and around the HP compositional field ($\text{K}_2\text{O}/\text{Na}_2\text{O} = 1.03\text{--}1.50$, $\text{CaO}/\text{Al}_2\text{O}_3 = 0.36\text{--}0.53$), except for one MI overlapping with the LP field ($\text{K}_2\text{O}/\text{Na}_2\text{O} = 0.66$, $\text{CaO}/\text{Al}_2\text{O}_3 = 0.66$). Melt embayments show comparable compositions of $\text{K}_2\text{O}/\text{Na}_2\text{O} = 1.06\text{--}1.24$ and $\text{CaO}/\text{Al}_2\text{O}_3 = 0.44\text{--}0.47$. Melt inclusions and embayments are Fe-rich ($\text{FeO} = 8.52\text{--}11.49$ wt.%), with values comparable to melt inclusions of 3 May 2009.

Melt inclusions of the 3 May 2009 event ($\text{CaO}/\text{Al}_2\text{O}_3 = 0.34\text{--}0.53$; $\text{K}_2\text{O}/\text{Na}_2\text{O} = 1.08\text{--}1.61$) also plot in the HP compositional field and are slightly less Fe-rich than the 24 November melt inclusions ($\text{FeO} = 8.12\text{--}11.08$ wt.%). Our studied glassy groundmasses of 3 May 2009 are compositionally more evolved than whole-rocks ($\text{MgO} = 6.07$ wt.%, $\text{CaO} = 11.36$ wt.%; La Felice and Landi 2011).

Olivine composition and Fe–Mg diffusion profiles

We selected olivine crystals mostly belonging to the 0.5–1 mm fraction size and few to the 250–500 μm fraction,

with morphologies varying from tabular to euhedral and no difference being evident between the two size fractions.

Two main populations (A and B) of olivine crystals were identified in the samples of the 3 major explosions:

- Group A comprises variably reverse-zoned olivine crystals that constitute the most abundant population (76% of the total); these can be further subdivided into subgroup A1, with more pronounced reversely-zoned olivine crystals with Fo_{69-71} cores and Fo_{76-83} rims, and subgroup A2 that are slightly reversely zoned olivine crystals with Fo_{69-71} cores and Fo_{72-73} rims. Olivine crystals belonging to these two subsets contain abundant melt inclusions mostly clustering in the core and in some cases have developed a variably thick (50–100 μm), Mg-rich reaction rim (Fig. 5c).

- Group B consists of olivine phenocrysts (24%) that are compositionally homogeneous (Fo_{70-71}) and bear a single, large melt inclusion (Fig. 6a).

Olivine crystals from the 8 November 2009 are mostly homogeneous with Fo_{70-72} (67%; group B), while 33% are slightly reversely zoned with Fo_{70-71} cores and Fo_{72-75} rims (subgroup A2) (Fig. 6a). Reaction rims are rare and limited to olivine crystals of subgroup A2; these are less developed compared to the 24 November 2009 olivine crystals. There is no correlation between MIs occurrence within the phenocrysts and MIs size. Olivine phenocrysts of 3 May 2009 are almost all homogeneous and show evolved compositions (Fo_{70-73} ; group B), suggesting they were inherited from the HP magma (Fig. 6a).

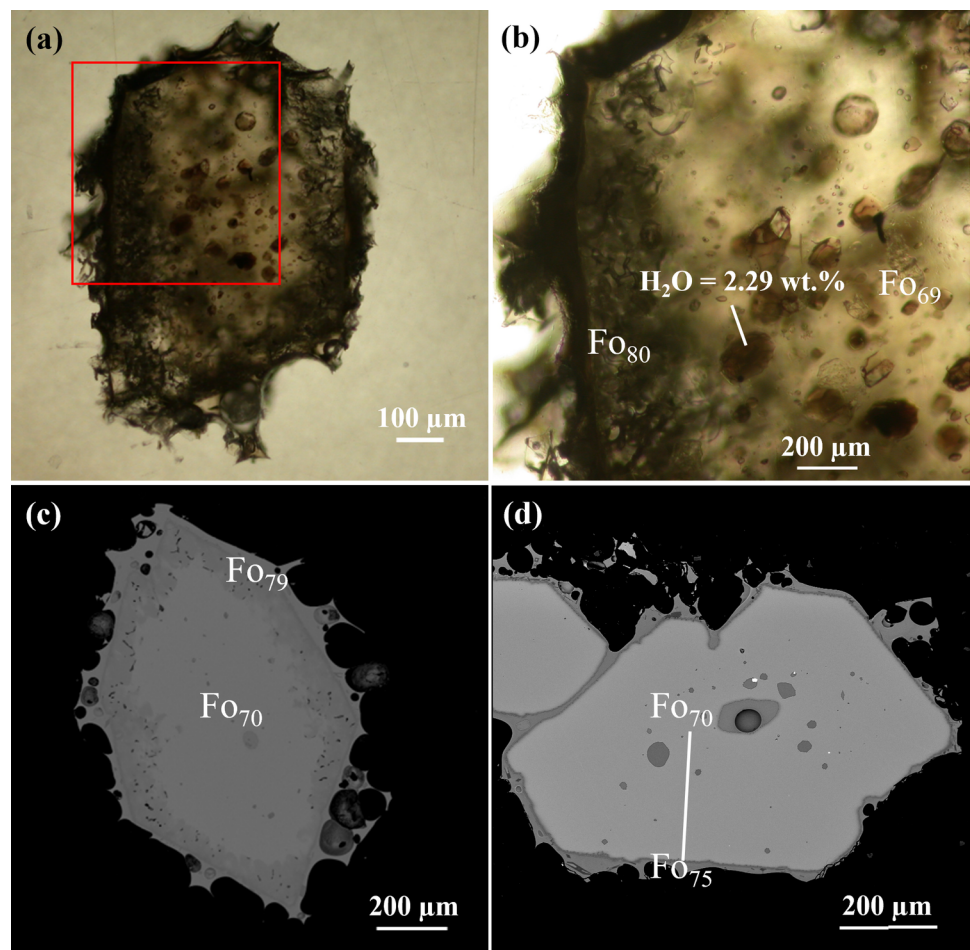
For the 24 November 2009 major explosion, Fe–Mg diffusion timescales range from a few days to 4 weeks (Fig. 6b), with only one sample (STN8-02) indicating timescales on the order of hours. Overall, these timescales are shorter than those inferred for the 19 July 2020 major explosion (~20–25 days, max. of 2 months; Voloschina et al. 2023), for the July and August 2019 small-scale paroxysms (5–115 days; Métrich et al. 2021) and the 1456 and 1930 CE large-scale paroxysms (2–58 days; Métrich et al. 2021). Diffusion profiles of the 8 November 2009 major explosion indicate timescales significantly shorter than those associated to other major explosions, in the order of only hours to a maximum of 1.5 day.

Olivine spot compositions and diffusion profiles are reported in Table S4 (in Supplementary Material 1) and Table S7–S9 (in Supplementary Material 2), respectively, while timescale results are reported in Figs. 6b and S11 (Supplementary Figures) and in Table S10 (Supplementary Material 2).

Volatiles in melt inclusions

Melt inclusions from the 24 November 2009 olivine crystals have H_2O contents that range from 0.22 to 2.37 wt.% (Fig. 7). Within this variability, melt inclusions with lower

Fig. 5 **a** Transmitted light microphotograph of an evolved olivine phenocryst from the 24 November 2009 clasts. Red insert is in **b** where is shown a high- H_2O melt inclusion in intermediate position between the evolved Fo_{69} core and the more primitive Fo_{80} olivine rim. This is a case of post-entrapment rehydration of melt inclusion by the LP melt observed in several reversely zoned olivine phenocrysts from the 24 November 2009. **c** Olivine phenocryst from 24 November with thick Mg-richer reaction rim and evolved resorbed core. **d** Fe–Mg diffusion profile measured in an olivine phenocryst from the 8 November hosting several large melt inclusions in its core. Numbers refer to forsterite component in olivine ($\text{Fo}\% = 100 \times \text{Mg}/(\text{Mg} + \text{Fe})$)



H_2O contents (0.22–0.87 wt.%) are hosted by evolved and homogeneous or slightly reverse Fo_{70} olivine. Melt inclusions with higher H_2O contents (1.42–2.37 wt.%) are hosted mostly in evolved olivine crystals (Fo_{69-71}) that are strongly reversely zoned (up to Fo_{83}), and only in one case slightly reversely zoned (Table S2 in Supplementary Material 1). Some of these H_2O -rich MIs occupy an intermediate position between the evolved core and the more primitive rim (Fig. 5a, b). CO_2 is always below the detection limit (50 ppm), except for one larger ($180 \times 100 \mu\text{m}$) and bubble-bearing melt inclusion — located close to the rim of a homogeneous Fo_{70} olivine — that contains 576 ppm of CO_2 . Sulphur and chlorine are highly variable, ranging between 60 and 993 ppm (with a primary mode at 309 ppm) (Fig. 8) and 660–2990 ppm (primary mode at 1585 ppm), respectively. Embayments have lower and less variable H_2O contents, ranging from 0.31 to 0.57 wt.%, CO_2 below the detection limit, and sulphur and chlorine ranging between 216–220 ppm and 1320–1180 ppm (values on one melt embayment), respectively.

Melt inclusions of the 8 November 2009 samples have H_2O contents from 0.55 wt.% to below the detection limit, and CO_2 always below the detection limit. S contents range from 55 to 777 ppm, with a primary mode at 187 ppm. Chlorine contents range between 1050 and 2600 ppm, overlapping the 24 November 2009 melt inclusion range. Melt embayments have H_2O contents of 0.18–0.47 wt.%, comparable to melt inclusions, and CO_2 below the detection limit. Sulphur range between 168 ppm to below the detection limit and chlorine contents vary between 1150 and 1310 ppm.

Melt inclusions of 3 May 2009 have H_2O contents ranging from 0.15 to 0.75 wt.%, CO_2 always below the detection limit, S ranging from 619 ppm to below the detection limit and Cl of 772–2779 ppm.

Sulphur and chlorine contents in glassy groundmass wetting the olivine phenocrysts range between 31–285 ppm and 910–1405 ppm, respectively. These values are lower than those of melt inclusions and are comparable for the 24 November, 8 November and 3 May 2009 samples.

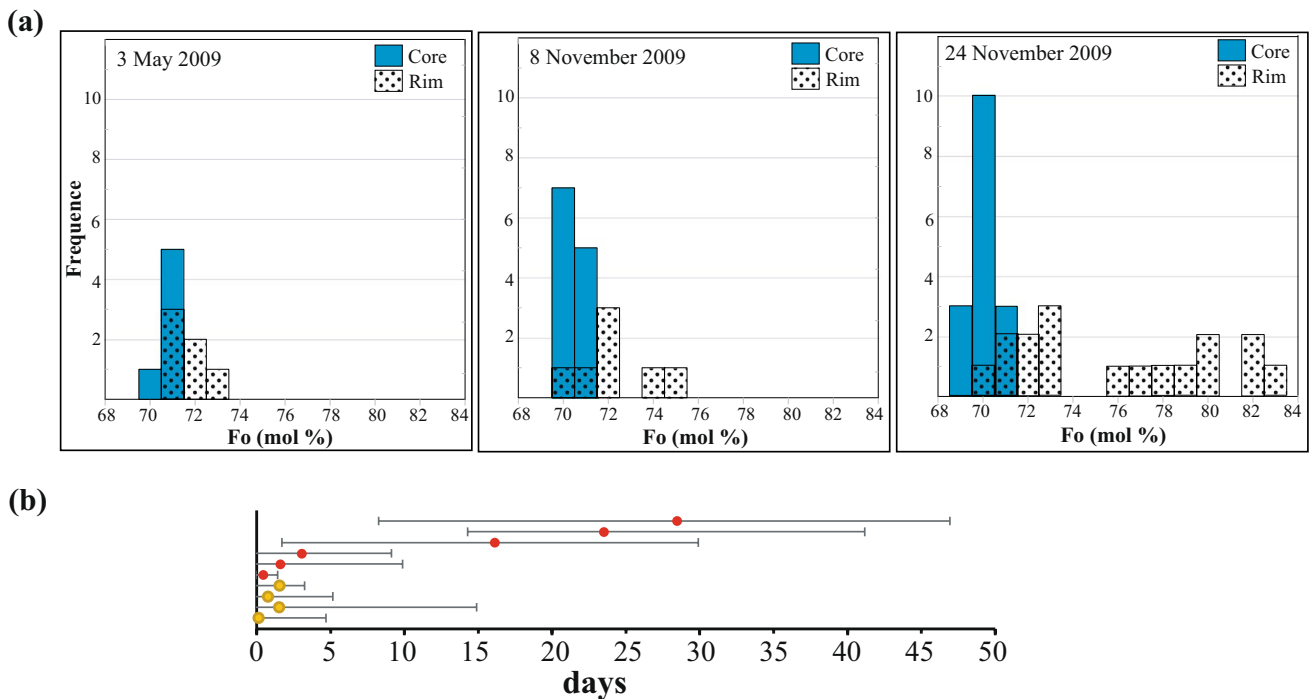


Fig. 6 **a** Histogram of olivine populations from the 24 November, 8 November and 3 May 2009 major explosions, comparing core and rim composition. **(b)** Diffusion timescales (in days) calculated in DIPRA (Girona and Costa 2013) in olivine phenocrysts from the 8

November (yellow filled dots) and 24 November (red filled dots) 2009 major explosions and associated uncertainty (based on the uncertainty in temperature and concentration) indicated with grey error bars

Volcanic gas composition and fluxes

The here presented melt inclusion results are complemented with previously published (Aiuppa et al. 2011) results for the composition/flux of the Stromboli volcanic gas plume. Results (Fig. 9) indicate that, throughout 2009, the SO_2 flux remained stable and low, with a mean average of 142 ± 59 (σ) tons/day. In contrast, the CO_2 flux was more variable and ranged between 65 and 7004 tons/day. The (molar) CO_2/SO_2 ratio varied between ~ 1 and 31.

Remarkable gas changes preceded the 24 November major explosion (Fig. 9). In particular, the explosion was associated with an abrupt increase of the CO_2/SO_2 ratio that started only a few hours before the explosive event. The CO_2 flux also abruptly increased to 2736 tons/day the day of the major explosion and peaked at 7004 tons/day the day after. The 3 May major explosion was also associated with some, less prominent gas changes. The median molar CO_2/SO_2 ratio in the volcanic plume (Fig. 9a) was ≤ 10 from January until late April, when it started to increase to > 15 molar ratio, remaining high until the 3 May 2009 major explosion. Few days before the 3 May major explosion, the CO_2 flux also increased, peaking at 1836 tons/day on 1 May. The 8 November explosion was not associated with any evident gas change, although it occurred during a phase of relatively high CO_2 fluxes.

Discussion

Pre-eruptive magma storage conditions

One critical question to address at Stromboli is whether (i) the shallow conduit system or (ii) the deep magma feeding system (or both) are involved in the generation of major explosions. Pre-eruptive volatile contents, as recorded by melt inclusions, can help to constrain the depth(s) of pre-eruptive magma storage (i.e., Anderson and Brown 1993; Métrich and Wallace 2008; Wallace et al. 2015b, 2021; Lerner et al. 2024).

Most melt inclusions of the 24 November 2009 display the typical composition of HP melt ($\text{K}_2\text{O} > 4$ wt.%; $\text{CaO}/\text{Al}_2\text{O}_3 \sim 0.5$), with S content as low as 0.1–0.2 wt.%, and CO_2 below the detection limit. Conversely, they display highly variable H_2O contents from 0.4 wt.% to 2.37 wt.% (Figs. S4, S5 in Supplementary Figures). Both relatively low H_2O (0.4 wt.%) and Al_2O_3 (~ 15 wt.%) concentrations are indicative of significant plagioclase crystallization typical of the crystal-rich magma residing in the upper parts of the volcano (Landi et al. 2004, 2022). Accordingly, these melt inclusions record a strongly degassed magma. Instead, relatively high H_2O content (> 1.5 wt.%,

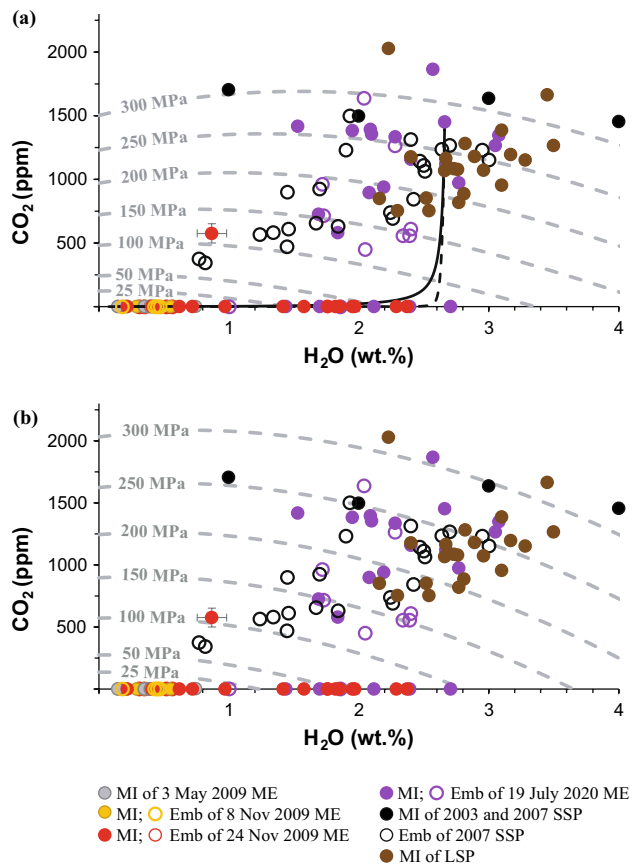


Fig. 7 Dissolved CO₂ and H₂O (wt.%) in melt inclusions (filled dots) and embayments (empty dots) from the 3 May, 8 November and 24 November 2009 major explosions (ME) in comparison with 19 July 2020 major explosion (Voloschina et al. 2023), 5 April 2003 and 15 March 2007 small-scale (SSP) paroxysms (Métrich et al. 2005, 2010) and historic large-scale (LSP) paroxysms (Bertagnini et al. 2003; Métrich et al. 2010). Isobars in MPa (grey dashed lines) were calculated in VESICAL (Iacovino et al. 2021) using an averaged melt composition from the 19 July 2020 major explosion (see Table S6, Supplementary Material 1) at 1150 °C. **a** Isobars and a general degassing path (solid line=open system; dashed line=closed system) of a 19 July 2020 melt inclusion were modelled using the MagmaSat solubility model (Ghiorso and Gualda 2015). **b** Isobars were modelled using the Iacono-Marziano solubility model (Iacono-Marziano et al. 2012). The maximum errors for H₂O and CO₂ (13%) determined in the 3 May, 8 November and 24 November 2009 melt inclusions and embayments are indicated as error bars

Fig. S5) is inconsistent with very low dissolved sulphur concentrations (Fig. 8), massive plagioclase crystallization (Di Carlo et al. 2006) and relatively low Al₂O₃ content which has been analysed (Table S2). The high-H₂O melt inclusions of the 24 November event are entrapped in strongly reversely-zoned olivine crystals from Fo₆₉₋₇₁ in the core to Fo₈₂₋₈₃ in the rims. These latter olivine compositions are in equilibrium with LP-type glassy groundmass (CaO/Al₂O₃ ~ 0.60, Fig. S3 in Supplementary Figures). As a whole, these observations strongly suggest H₂O (H⁺)

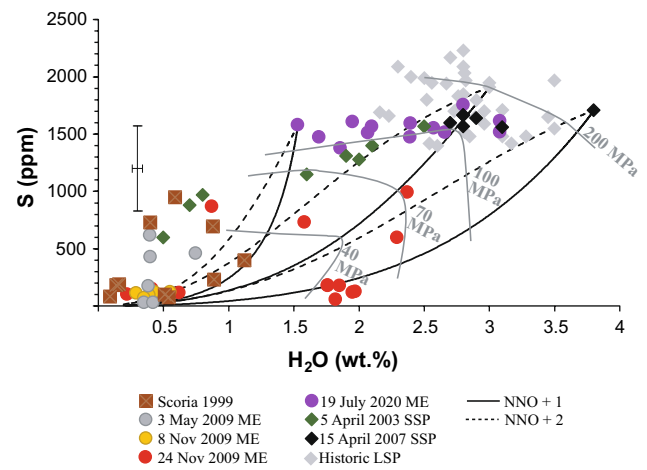


Fig. 8 Sulphur (ppm) vs H₂O (wt.%) contents in melt inclusions of the 3 May, 8 November and 24 November 2009 major explosions (ME; this study), in comparison with the 19 July 2020 major explosion (Voloschina et al. 2023), 5 April 2003 small-scale paroxysm (SSP; Métrich et al. 2005), 15 March 2007 small-scale paroxysm (SSP; Métrich et al. 2010) and historic large-scale paroxysms (LSP; Bertagnini et al. 2003; Métrich et al. 2010). Black lines show a general degassing path modelled by using the Iacono-Marziano model (Iacono-Marziano et al. 2012). Runs were performed at 1150 °C, assuming a fo₂ NNO+1 (black solid line) and NNO+2 (black dashed line). Grey lines indicate isobars in MPa. Average error relative to S=31% and maximum error for H₂O=13% are indicated as error bars

re-equilibration of the initially almost dry melt inclusions through the olivine (Fo₆₉₋₇₁) as experimentally reported for the olivine-hosted MIs of the Galapagos Plateau basalt (Portnyagin et al. 2008). In all melt inclusions, CO₂ is below the detection limit, which suggests extensive degassing took place before MI entrapment. A maximum CO₂ content of 576 ppm has been detected exclusively in an intermediate and bubble-bearing melt inclusion (sample STN8-15-M7). The measured value represents only the CO₂ content dissolved in the melt; however, CO₂ is known to be severely partitioned into shrinkage bubble (Hartley et al. 2014; Moore et al. 2015; Wallace et al. 2015a, b, 2021; MacLennan 2017; Robidoux et al. 2018; Tucker et al. 2019; Rasmussen et al. 2020; Wieser et al. 2021; Buso et al. 2022). As such, the measured CO₂ content likely underrepresents the total amount originally dissolved in the melt. This melt inclusion is located near the Fo₇₀ olivine rim which is in equilibrium with the groundmass (HP), hence was not affected by post-entrapment processes of rehydration, consistent with its low H₂O content (0.87 wt.%).

We exclude from the P_{sat} calculations those melt inclusions that recorded high-H₂O content in the 24 November 2009 event, as they have been affected by secondary rehydration processes, and therefore their values do not represent pressure values at the time of the MI entrapment. Similarly,

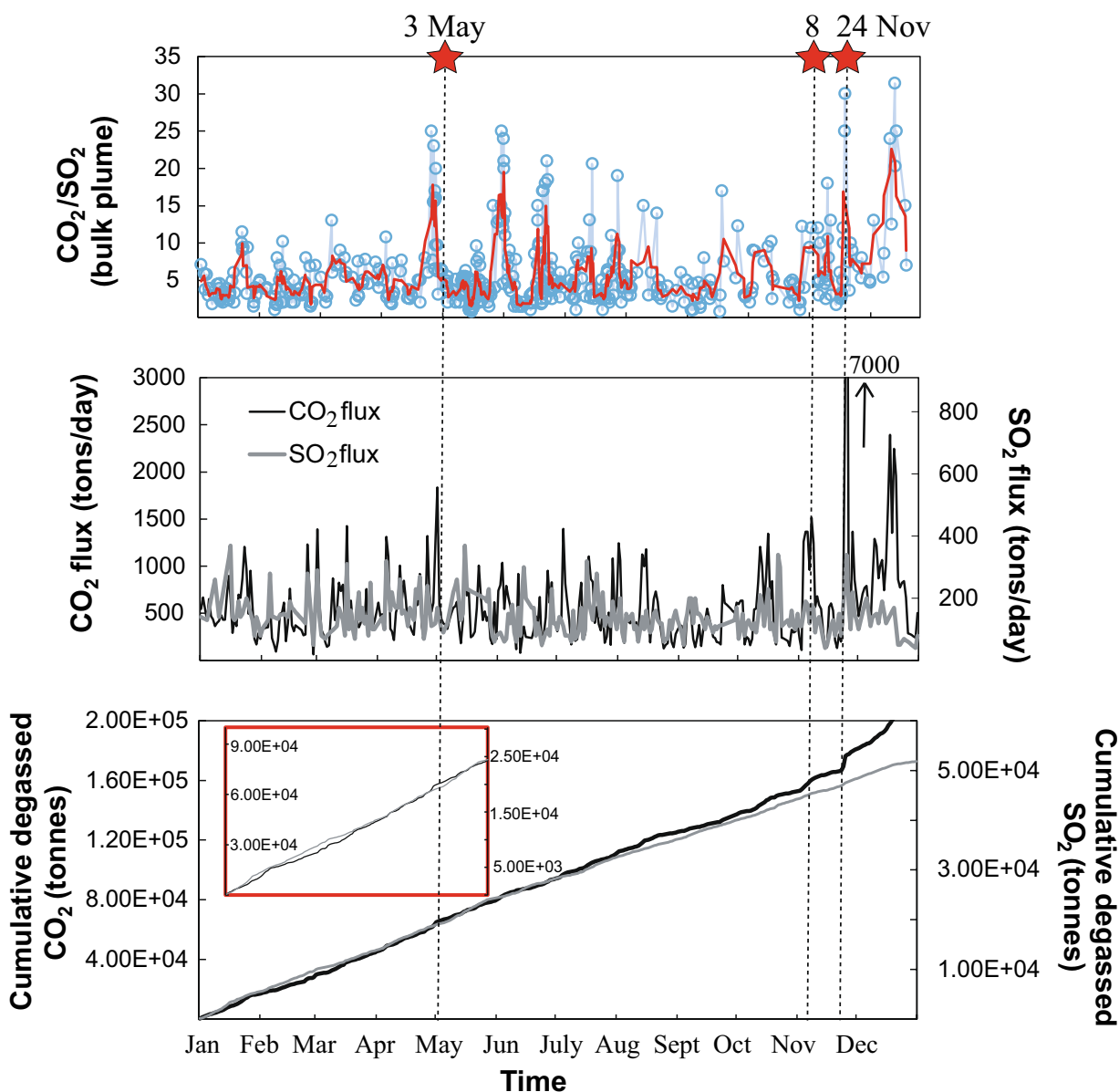


Fig. 9 Gas plume observation at Stromboli during the 2009 year. **a** CO₂/SO₂ ratio in the bulk plume. **b** CO₂ (black line) and SO₂ (grey line) fluxes (in tons/day). **c** Cumulative CO₂ (black line) and SO₂ (grey line) fluxes. The CO₂ scale is 3.3 times higher than the SO₂ scale in order to normalise it to the time-averaged CO₂/SO₂ ratio of Stromboli’s emission during the observation period. The red line

indicates the weekly mobile average, whereas the red stars indicate the onset of the 3 May, 8 and 24 November major explosions. The red insert shows a focus on the 3 May 2009 event: the CO₂ flux decelerated relative to the SO₂ flux in the months prior the onset of the eruption, followed by a convergence of the two cumulative CO₂ and SO₂ fluxes

based on the position close to the olivine’s rim and intermediate-composition of STN8-15-M7, we cannot rule out that it recorded a late episode of magma mixing, thus its volatile composition was excluded from further calculation.

Here, our saturation pressure calculations are based on a comparison between two models, the MagmaSat solubility model (Ghiorso and Gualda 2015) and the Iacono-Marziano solubility model (Iacono-Marziano et al. 2012), both implemented in VESIcal (Iacovino et al. 2021). The former has been calibrated for a wide range of compositions,

temperatures and pressures and has been used for the pressure calculations based on high H₂O and CO₂ contents in melt inclusions for the 19 July 2020 major explosion (H₂O = 1.53–3.64 wt.% and CO₂ = 580–1866 ppm; Voloschina et al. 2023) as well as the recalculation of MIs for the 2003 and 2007 paroxysms (Métrich et al. 2010). On the other hand, while the calibration range of the experiments that underlie the Iacono-Marziano solubility model is more limited, it is specifically calibrated on alkali-basaltic melt compositions (Iacono-Marziano et al. 2012). Most

importantly, the latter can be implemented in the Sulfur_X model of Ding et al. (2023), thus allowing to model sulphur degassing (see following sections).

The H₂O vs. CO₂ plot (Fig. 7a, b) shows the isobars modelled at 1150 °C using an averaged LP intermediate melt composition, in equilibrium with Fo₈₀₋₈₃ olivine crystals, taken from the 19 July 2020 dataset (Voloschina et al. 2023; composition resumed in Table S6, Supplementary Material 1), comparing the isobars modelled using the MagmaSat (Fig. 7a) or the Iacono-Marziano solubility model (Fig. 7b). What clearly emerges from this comparison between the two plots (Fig. 7a, b) is that the differences arising from the use of these two different solubility models are minimum at low pressure values (2009 major explosions), while differences are higher at high pressure values (19 July 2020 and paroxysms), showing deviation on saturation pressures up to 30 MPa for the 19 July 2020 major explosion.

Consequently, for the low H₂O and CO₂ contents measured in melt inclusions of all the 2009 major explosions, both models yield similarly low entrapment pressures, lower than 10 MPa (Iacono-Marziano et al. 2012 solubility model) and lower than 7 MPa (MagmaSat solubility model). These pressures correspond to a shallow inferred depth, lower than 1 km below the sea level (b.s.l.) (assuming an average crustal density of 2700 kg/m³).

To complement the above set of information, and in particular considering that saturation pressures for 2009 melt inclusions are based on H₂O contents only, we also attempt a degassing estimate based on the sulphur (S) vs. H₂O systematics (Fig. 8). We used Sulfur_X v1.1 (Ding et al. 2023) to model the S degassing behaviour (in combination with H₂O and CO₂) during magma ascent and decompression (down to 1 atm) under closed-system conditions. This code combines a calibration of experimentally derived sulphur partition coefficients between silicate melt and co-existing vapor phase with an existing C-O-H solubility model (in this case, the Iacono-Marziano solubility model). However, large uncertainties arise for sulphur partition coefficients at < 25 MPa, as no experimental sulphur partition coefficients exist at such low-pressure conditions.

Three couples of independent model degassing paths were generated, all at 1150 °C and fO₂ ranging from NNO + 1 (black solid line) to more oxidised (NNO + 2; black dashed line) conditions. Each of the three couples were obtained by initialising the model runs for a given set of starting melt compositions (composition reported in Table S6, Supplementary Material 1), and exemplified by MIs of the: (i) 19 July 2020 major explosion (H₂O = 1.5 wt.% and S = 1582 ppm; Voloschina et al. 2023); (ii) 15 March 2007 small-scale paroxysm (H₂O = 3.8 wt.% and S = 1710 ppm; Métrich et al. 2010) and (iii) a historic large-scale paroxysm (H₂O = 3 wt.% and S = 1910 ppm; Métrich et al. 2010) with high S and H₂O contents.

Results of model simulations of degassing are compared against MIs information in Fig. 8. Model simulations indicate sulphur begins to exsolve at rather high pressures, consistent with experimental data on Stromboli oxidised melts (Lesne et al. 2011) and previous melt inclusions study (Métrich et al. 2021), indicating S degassing from as deep as at 150 MPa (~ 5 km b.s.l.). Below this pressure, model lines reasonably reproduce the coupled drop of S and H₂O concentrations recorded in MIs of the 2009 major explosions (this study), and also of glass embayments of the 2020 major explosion and of the 2003 paroxysms. This comparison therefore suggests shallow entrapment pressures for the 2009 melt inclusions, that are consistent with those previously calculated from H₂O contents only (entrapment pressures of < 10 MPa; Fig. 7). By contrast, in the high-pressure range (150–200 MPa), MIs from the 19 July 2020 major explosion and from the small- and large-scale historic paroxysms exhibit an overall tendency of decreasing H₂O at constant S, interpreted to represent isobaric magma dehydration caused by flushing by CO₂-rich bubbles (Métrich et al. 2010; Voloschina et al. 2023).

In summary, the inferred MIs entrapment depths indicate that the 2009 major explosions predominantly drained material originally stored in the upper feeding conduit system, at source depths lower than 400 m b.s.l., corresponding to the shallower portion of the dyke-conduit feeding system, which extends from surface to ~ 1–2 km b.s.l. depth (Chouet et al. 2003, 2008; Patané et al. 2017).

Pre-eruptive processes and mixing to eruption timescales

During the last two decades, extensive scientific research has corroborated the idea for the existence of a vertically extended plumbing system at Stromboli, in which a complex interplay of fractional crystallization, crystal dissolution, magma mixing and degassing concur to control magma evolution (Bertagnini et al. 2003; Francalanci et al. 1988, 1989, 2004; Landi et al. 2004, 2008, 2022; Métrich et al. 2001, 2010, 2021). By integrating compositional results for host olivine crystals, melt inclusions and glassy groundmass, we here attempt to refine our understanding of the driving pre-eruptive processes of the intermediate, yet highly hazardous category of explosive events represented by the major explosions.

One key observation is that olivine crystals in all the 2009 products are relatively low in Mg, with evolved (Fo₆₉₋₇₂) olivine cores, and relatively flat Fe–Mg compositional profiles over hundreds of micrometres (Fig. S11 in Supplementary Figures). This, combined with the dominant HP compositional affinity of the melt inclusions (Figs. 3, 4), and their low volatile contents, suggest these minerals formed during protracted magma storage in a shallow reservoir. If then

major explosions are triggered by overpressure development in the shallow HP magma reservoir, the question that arises is what drives this overpressure, and over what timescales.

The rapid ascent of deeply-stored LP magma, and its mingling with the shallow HP magma, has long been invoked as a key driver of the larger than normal explosions at Stromboli (Bertagnini et al. 2003; Di Carlo et al. 2006; Métrich et al. 2001, 2010, 2021; Pichavant et al. 2009, 2022). The recurrent reverse zoning (Fig. 6) observed in olivine phenocrysts from the 24 November major explosion (with rims up to Fe_{83}) and (to a minor extent) from the 8 November major explosion, supports pre-eruptive interaction between shallow-stored HP phenocrysts and Mg-richer melts. Two additional lines of evidence corroborate this process. First, for both eruptions, the glassy groundmass is compositionally more primitive than the olivine-hosted MIs, partly overlapping the LP compositional field (Figs. 3, 4). Secondly, the 24 November olivine phenocrysts contain abundant (and pervasive) resorption textures (Fig. 5a–c, Fig. S1 in Supplementary Figures; these are also present — although less commonly — in the 8 November olivine crystals). These resorption textures implicate that, following a long-lived period of equilibrium crystallization within the shallow resident magma (during which cores with evolved composition $\sim Fe_{70}$ were formed), higher-temperature and water-richer melts rapidly entrained the system, causing the redissolution of the phenocryst rims (Streck 2008). Taken together, these lines of evidence suggest a trigger mechanism for major explosions in which deeply rising, volatile-rich LP magma is rapidly injected into the shallow HP magma storage zone (see also Voloschina et al. 2023). The presence of resorbed olivine crystals is fully consistent with the reactivation — during the more energetic major explosions (particularly the 24 November event) — of the stagnant crystal-mush zone by the influx of the deep LP magma (Bertagnini et al. 2003; Di Stefano et al. 2020; Francalanci et al. 2004, 2005, 2012; Métrich et al. 2021; Petrone et al. 2022).

The timescales of this pre-eruptive HP-LP interaction can be inferred from modelling of diffusion profiles in olivine rims (e.g., Costa et al. 2020) (Fig. 6b). Overall, olivine cores of the November major explosions suggest a protracted storage in the shallow reservoir, whereby olivine crystals completely equilibrate through diffusion in steady-state conditions. In particular, olivine crystals of the 8 November explosion document short mixing to eruption timescales, in the order of hours to < 2 days. These temporal timescales are shorter than those required for extensive mixing between HP and LP magmas — as, at least in principle, required to explain the intermediate compositions observed in the glassy groundmasses from the 8 November eruption (La Felice and Landi 2011; Landi et al. 2022). We hence propose rapid HP-LP interaction was favoured (Perugini et al. 2003; De

Campos et al. 2011) by vigorous ascent of deeply-rising gas bubbles, as suggested by gas data (see below).

A wider range of timescales, from hours to a maximum of 4 weeks, is recorded by olivine crystals of the 24 November explosions. The 4 weeks upper limit is longer than the eruptive inter-event time (the 16 days elapsed between the 8 and 24 November explosions). We argue that degassed and dense (unerupted) HP magma, left from the previous eruptive activity in the shallow reservoir, was re-mobilised by the ascending bubbly LP magma in the eruption run-up.

In contrast to the November events, the investigated 3 May 2009 products show no evidence of more primitive (LP magma-like) glassy groundmass (Figs. 3–4; note that some rare, more primitive groundmass has been however reported for this eruption; Corsaro and Miraglia 2009a; La Felice and Landi 2011; Pioli et al. 2014, who analysed the distal fraction of the tephra deposit made by a few scattered lapilli). This absence/rarity of LP magma-like groundmass is consistent with the smaller LP erupted mass in the 3 May eruption ($1.8\text{--}2.9 \times 10^4 \text{ m}^3$, relative to $4.4\text{--}6.5 \times 10^5 \text{ m}^3$ for the 24 November eruption; Pioli et al. 2014). The 3 May major explosion olivine crystals are also compositionally homogeneous (only one olivine exhibits some slight reverse zoning). In summary, these results suggest that the LP magma was only marginally involved in the triggering of the 3 May major explosion, requiring a distinct mechanism.

Clues from volcanic gas plume monitoring

Volcanic gas observations corroborate a distinct trigger mechanism for the May and November 2009 major explosions. We rely here on the gas dataset already presented and discussed in Aiuppa et al. (2011) (Fig. 9a, b). These data are used to calculate the cumulative masses (in tonnes) of degassed CO_2 and SO_2 throughout 2009, illustrated in Fig. 9c (note that the CO_2 scale is 3.3 times higher than the SO_2 scale in order to match the time-averaged CO_2/SO_2 mass ratio). This diagram highlights a contrasting degassing behaviour for the periods preceding the November and May 2009 explosions, respectively.

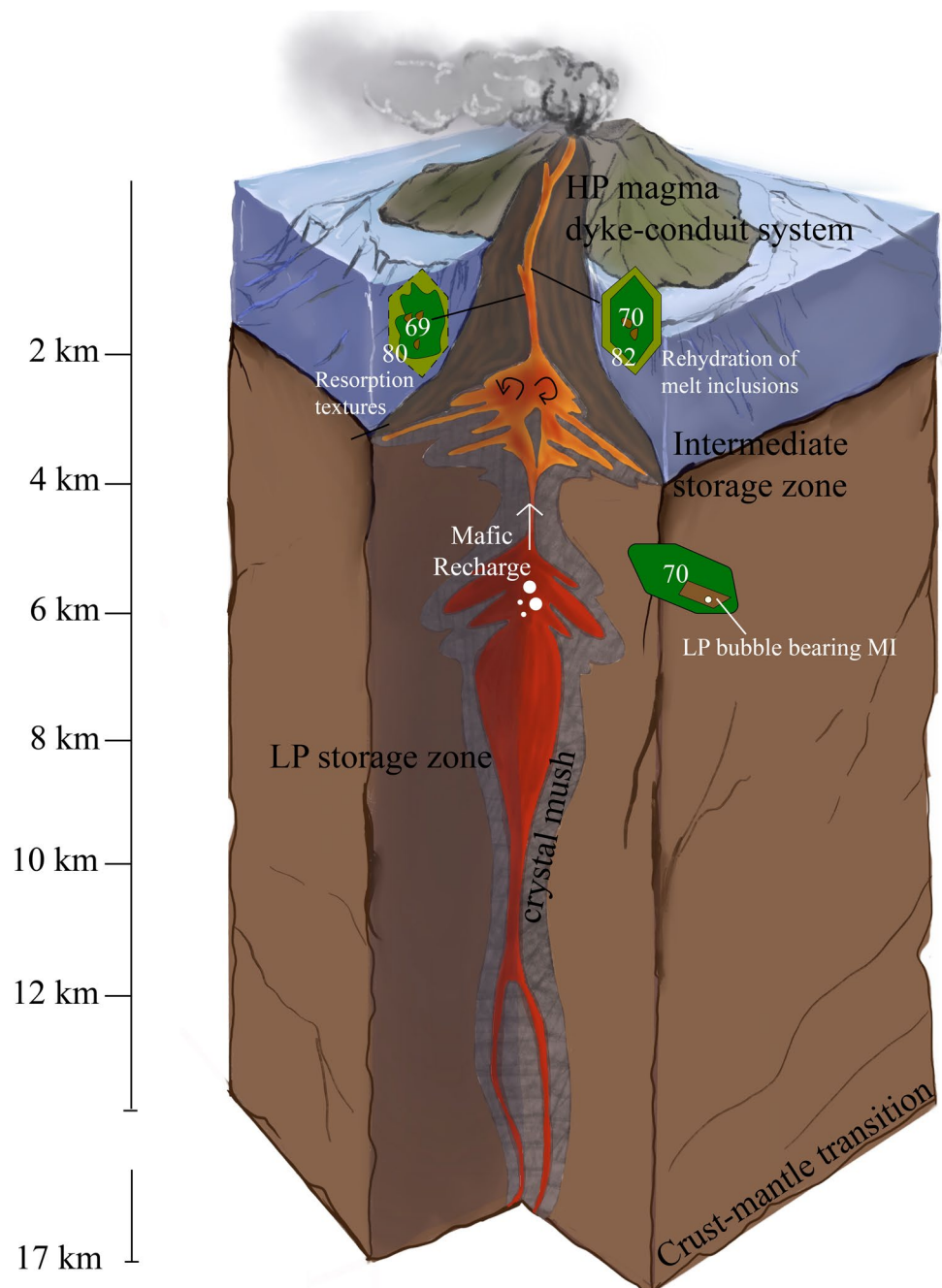
In the months prior to the November explosions, the CO_2 flux accelerated relative to the SO_2 flux, as indicated by the diverging cumulative trends (Fig. 9c). This CO_2 acceleration started sometime in late summer 2009, and became more pronounced in mid- to late-October, e.g., weeks to days prior to the first major explosion on November 8 (Fig. 9c). The peak of CO_2 flux was observed in correspondence of the 24 November explosion (see Fig. 9b). Taken together, and considering the CO_2 -rich nature of Stromboli's deep LP magma (relative to the HP magma that is CO_2 -depleted), these data are consistent with a mechanism in which the November explosions were preceded by an anomalous inflow of CO_2 -rich gas bubbles, transported by (with) the ascending LP

magma (as testified by olivine zoning and groundmass chemistry; see the “Pre-eruptive processes and mixing to eruption timescales” section). A similar mechanism was proposed for the July 2020 major explosion (Voloschina et al. 2023) and is hence likely to be a common characteristic of the more energetic events (those erupting larger LP pumice volumes).

No obvious CO₂ acceleration was observed prior to the 3 May 2009 event. In contrast, in the months prior the 3 May major explosion, the CO₂ flux decelerated relative to the SO₂ flux (from February to mid-March, see inset in Fig. 9c), followed by a recovery phase during which the

two cumulate flux trends converged again (e.g., the CO₂/SO₂ mass ratio returned to its time-averaged value). The peak CO₂ was observed in the days prior to the 3 May blast. Aiuppa et al. (2011) explained these cycles of decelerating/accelerating CO₂ to be caused by (i) an initial phase of gas bubbles retention at depth (causing a deceleration of the CO₂ flux), gradually accumulating a foam at some (rheological or structural) discontinuity (Allard 2010; Aiuppa et al. 2011; Caricchi et al. 2024), followed by (ii) a precursory phase of passive gas leakage from the growing foam (CO₂ flux acceleration; Phillips and Woods 2001), and subsequent (iii)

Fig. 10 Interpretative model of Stromboli’s plumbing system explaining the pre-eruptive dynamics of the 24 November 2009 major explosion, based on (i) major element and sulphur contents on melt inclusions, embayments, glassy groundmass and host olivine crystals, (ii) dissolved H₂O and CO₂ contents on melt inclusions and embayments, and (iii) olivine textures



catastrophic collapse of the foam, leading to the fast ascent and surface bursting of gas slugs during the major explosion (Allard 2010). This mechanism does not require any important pre-event upward migration of LP magma, which would be difficult to reconcile with our mineral and glass chemistry results (see above).

In summary, gas and petrological results (host olivine composition, MIs and glassy groundmass compositions) indicate there may be no single mechanism to explain the generation of major explosions. Rather, these events can either be gas-driven (e.g., caused by gas accumulation and then failure of a foam) or magma-driven (e.g., caused by the upward migration of small pockets of deep volatile-rich magma, causing pressure build-up in the shallow HP reservoir). In this interpretation, key factors in modulating the diversity of explosion magnitude are (i) the relative proportions of gas and LP magma involved in the process, and (ii) the depth at which the gas bubble retention event (in the gas-driven scenario), or the LP-HP interaction event (in the LP magma-driven scenario) occur (Allard 2010; Métrich et al. 2021; Landi et al. 2022). We conclude that the low-intensity, 3 May 2009 major explosion was critically driven by the gas accumulation within the shallower (< 1 km b.s.l.) magma reservoir, implicating a marginal involvement of the deep LP plumbing system. In contrast, a more active role for the deep LP magma is implicated for both November 2009 events. In this case, the rising LP magma interacted (over timescales of days to weeks) with the shallow HP magma reservoir and with the surrounding crystal mush zone, ultimately causing a more deeply-sourced, and hence more energetic, series of explosions (Andronico and Pistolesi 2010) (Fig. 10).

Linking ground deformation patterns to eruptive intensity

Following our discussions of volcanic gas and petrological insights, we now focus on the analysis of the ground deformation measured during the 2009 major explosions.

In Fig. 11a, the ground deformation (tilt) filtered for the Very-Long Period (VLP) seismic signal is reported. The 3 May 2009 produced a ground tilt of ~0.2 μrad recorded by tiltmeters, while the 8 and 24 November events produced a ground tilt of 0.48 and 0.32 μrad , respectively. The measurements of the ground deformations confirm the 3 May 2009 as the lowest-intensity major explosion among the studied 2009 events, but also raise the question as to how the 8 November major explosion produced a higher ground deformation, despite having ejected smaller magma volumes and having smaller dispersal areas (10,000 m^2) than the 24 November event (15,000 m^2 ; Andronico and Pistolesi 2010). To explain this apparent discrepancy in the deformation amplitude, we highlight the 24 November is the result of two different blasts occurring ~15 s apart which contributed to a slightly larger total volume. In the whole, the amplitude

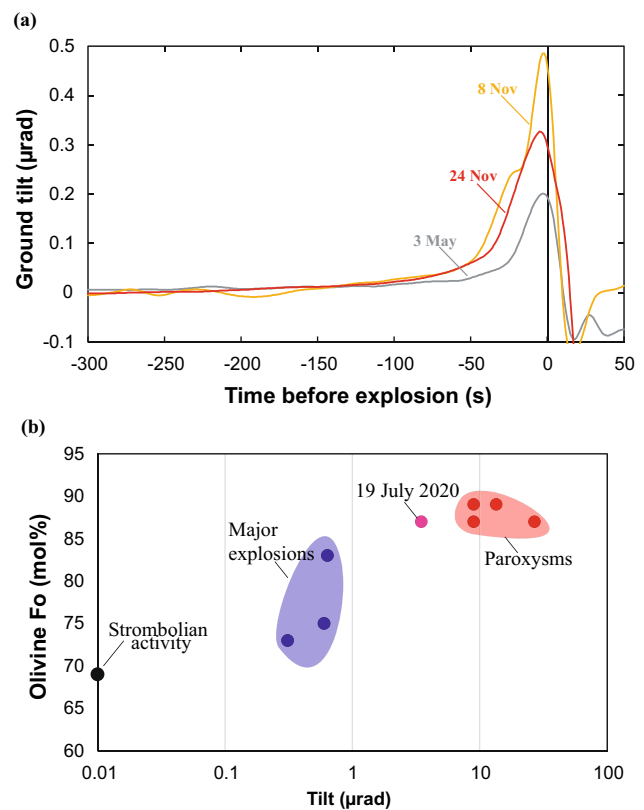


Fig. 11 **a** Ground deformation (tilt), filtered for the Very-Long Period signals, associated with the 2009 major explosions. Note that the chosen time frame (X axis) is consistent with the observation that the ground inflation becomes more visible ~300 s before the onset of each major explosion. **b** Semilog plot of tilt (μrad ; contaminated with VLP) against olivine composition (Fo mol%) of regular Strombolian explosions to major explosions (3 May, 8 November and 24 November 2009, 19 July 2020) and paroxysms at Stromboli (5 April 2003; 15 March 2007; 3 July and 28 August 2019)

of the pre-explosion deformation falls within the range typical of major explosions class and scales with the magnitude/intensity of each explosion, depending on the volumes of the ejected material, as previously found by Ripepe et al. (2021).

Lastly, if we correlate the ground deformation measured by tiltmeters during the Stromboli's eruptive activity, spanning from its regular Strombolian explosions to paroxysms, to their respective olivine (forsterite) compositions (Fig. 11b), it becomes evident that the ordinary activity is associated to low ground deformation (0.1 μrad) and evolved Fo_{70} olivine, while major explosions range instead from 0.31 to 0.64 μrad with olivine composition from Fo_{73} to Fo_{83} . An exception to this is the 19 July 2020 event (3.5 μrad ; Fo_{87} olivine), which falls between the 2009 events and the paroxysmal activity (9–27 μrad ; Fo_{87-89} olivine). What clearly emerges is also that, in agreement with the previous findings of Métrich et al. (2021) and Voloschina et al. (2023), the olivine chemistry display a good correlation with the tilt measurements, thus with the eruption intensity/magnitude of the explosions.

Conclusions

Stromboli exhibits eruptions ranging in size from regular Strombolian explosions to paroxysms, whose diversity is governed by the (variable) interplay between the shallow HP magma and gas \pm magma supply from depth (Pioli et al. 2014; Métrich et al. 2021; Voloschina et al. 2023). The regular Strombolian-type explosions are sustained by the shallow-stored HP magma (with rare LP clasts interpreted as fragments dragged to the surface by deep CO₂-rich gas bubbles; D’Oriano et al. 2011). In contrast, the roles of deeply sourced bubbles and LP magma increase with increasing eruption magnitude/intensity. For example, the smallest major explosions (at the upper boundary of regular Strombolian-type explosions) do not emit LP products (i.e., 8 September 1998, Bertagnini et al. 1999; 7 September 2008; Calvari et al. 2012), or they do in negligible amounts (3 May 2009, Corsaro and Miraglia 2009a; La Felice and Landi 2011; Pioli et al. 2014). In contrast, higher volumes of LP magma are emitted during the most violent major explosions (8 and 24 November 2009, Corsaro and Miraglia 2009b,c; Pioli et al. 2014; La Felice and Landi 2011; Landi et al. 2022; this study; 19 July 2020, Voloschina et al. 2023) and small to large-scale paroxysms (5 April 2003, 15 March 2007 and historic events, Bertagnini et al. 2003; Métrich et al. 2005, 2010), implicating the involvement of the deep plumbing system.

The results presented here indicate that the diversity in style and magnitude observed within the eruptive class of major explosions likely reflects distinct eruption triggering mechanisms. We have shown that the November explosions contain mineralogical, textural and petrological evidence for the injection of variable volumes of LP magma within the HP magma reservoir. This process is likely to have taken place over timescales of hours (8 November) to weeks (24 November), and to have caused overpressure development at the HP magma reservoir. In contrast, the 3 May products contain only homogeneous (un-zoned) olivine crystals with composition comparable to that of ordinary activity (\sim Fo₇₀). The lack of resorption textures and low volatiles in MIs, coupled with the small amount of LP glassy groundmass (Corsaro and Miraglia 2009a; La Felice and Landi 2011; Pioli et al. 2014), confirm that the LP magma ascent played only a marginal role. Rather, in combination with volcanic gas plume results, our results indicate the 3 May eruption may have been triggered by a pressurization of the shallow (< 1 km b.s.l.) HP magma reservoir, caused by the accumulation of gas bubbles at some (rheological or structural) discontinuity. These diverse causal mechanisms, combined with short incubation timescales (hours to weeks), may explain why major explosions have remained so difficult to forecast (Aiuppa et al. 2025).

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Author contribution Laura Insinga: writing — original draft preparation, writing — review and editing, methodology, formal analysis and investigation, data curation, visualization, conceptualization. Marija Voloschina: writing — review and editing, methodology, formal analysis and investigation, data curation, visualization, conceptualization. Paola Marianelli: writing — review and editing, methodology, formal analysis, investigation, data curation, visualization, validation, conceptualization. Erika Bartolomeo: formal analysis and investigation, data curation. Antonella Bertagnini: writing — review and editing, validation, conceptualization. Nicole Métrich: writing — review and editing, validation, conceptualization. Silvio G. Rotolo: writing — review and editing, methodology, investigation, data curation, visualization, validation, conceptualization, project administration, supervision. Alessandro Aiuppa: writing — review and editing, resources, methodology, formal analysis and investigation, data curation, visualization, validation, conceptualization, funding acquisition, project administration, supervision. Maurizio Ripepe: methodology, formal analysis and investigation, data curation, conceptualization. Marco Pistolesi: writing — review and editing, resources, methodology, investigation, data curation, visualization, conceptualization, project administration, supervision.

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Data availability All the data discussed in the manuscript are presented in the figures and in the supplementary material.

Declarations

Conflict of interests The authors declare no competing interests.

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